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OM protein - protein search, using sw model

Run on: January 27, 2005, 17:32:04 : Search time 86.5 Seconds
(without alignments)
116.120 Million cell updates/sec

Title: US-10-608-584-1
28
Perfect score: 1 YMFVFLVFLGSPFLINLILAVVAVMAY 28
Sequence:

Scoring table: OLIGO
Gapop 60.0 , Gapext 60.0

Searched: 2002273 seqs, 358729299 residues

Word size : 0

Total number of hits satisfying chosen parameters: 2002273

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 100 summaries

Database : A_Geneseq_23Sep04:*

1: geneeqp1980s:*
2: geneeqp1990s:*
3: geneeqp2000s:*
4: geneeqp2001s:*
5: geneeqp2002s:*
6: geneeqp2003as:*
7: geneeqp2003bs:*
8: geneeqp2004s:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	28	100.0	447	7	ADB78592 Human sod
2	28	100.0	1381	5	AAE20513 Human ion
3	28	100.0	1387	5	AAE20514 Human ion
4	28	100.0	1392	5	AAE20518 Human ion
5	28	100.0	1398	5	AAE20519 Human ion
6	28	100.0	1442	5	AAE20512 Human ion
7	28	100.0	1453	5	AAE20517 Human ion
8	28	100.0	1795	5	ADB78596 Human sod
9	28	100.0	1855	7	ADB78597 Human sod
10	28	100.0	1962	5	AAE20511 Human ion
11	28	100.0	1973	5	AAE20516 Human ion
12	28	100.0	1981	7	ABR83185 Human SCN
13	28	100.0	1998	5	AAE20510 Human ion
14	28	100.0	1998	5	ABR83184 Human SCN
15	28	100.0	1999	5	ABR806026 Human sod
16	28	100.0	2005	4	AAAB99676 Human adu
17	28	100.0	2005	4	AAAB99677 Human neo
18	28	100.0	2005	5	ABR83627 Human GEF
19	28	100.0	2005	7	ADB78604 Human sod
20	28	100.0	2005	7	ADB78603 Human sod
21	28	100.0	2005	7	ADB78605 Human sod
22	28	100.0	2005	7	ADC46947 Human SCN
23	28	100.0	2009	4	AAAB99674 Human adu
24	28	100.0	2009	5	AAE20515 Human ion
25	28	100.0	2009	5	ABG69292 Human sod

26	28	100.0	2009	5	ABG69291 Human sod
27	28	100.0	2009	5	ABG69293 Human sod
28	28	100.0	2009	5	ABG69289 Human sod
29	28	100.0	2009	5	ABG69290 Human sod
30	28	100.0	2009	5	ABR83626 Human GEF
31	28	100.0	2009	5	AAE16776 Human tra
32	28	100.0	2009	7	ADB78599 Human sod
33	28	100.0	2009	7	ADB78595 Human sod
34	28	100.0	2009	7	ADB78593 Human sod
35	28	100.0	2009	7	ADB78594 Human sod
36	28	100.0	2009	7	ADB78598 Human sod
37	28	100.0	2009	7	ABR83180 Human SCN
38	28	100.0	2009	7	ADE57563 Human pro
39	28	100.0	2009	7	ADE57561 Rat Prote
40	28	75.0	1835	2	AAE92316 Periphra
41	21	75.0	1977	2	AAE92317 Periphra
42	21	75.0	1977	8	ADL13028 Human pro
43	21	75.0	1978	7	ADL13028 Human ste
44	21	75.0	1978	7	ADL13028 Human pro
45	21	75.0	1978	7	ADL13028 Human pro
46	21	75.0	1984	2	AAE99639 Periphra
47	21	75.0	1984	2	AAE99639 Periphra
48	21	75.0	1984	7	ADE54547 Rat Prote
49	21	75.0	1984	7	ADE54551 Rat Prote
50	21	75.0	1984	7	ADE54553 Rat Prote
51	21	75.0	1989	2	AAE99640 Periphra
52	20	71.4	1836	7	ADE57388 Human pro
53	20	71.4	1836	7	ADE59630 Human pro
54	20	71.4	1836	7	ADE63029 Human pro
55	20	71.4	1836	8	ADQ17412 Human sod
56	16	57.1	1024	5	ABR804858 LDL recep
57	16	57.1	1950	7	ADB78607 Human sod
58	16	57.1	1951	4	AAE99678 Human adu
59	16	57.1	1951	4	AAE99679 Human neo
60	16	57.1	1951	7	ADE59628 Rat Prote
61	16	57.1	1951	8	ADL06576 Human tnm
62	16	57.1	1956	4	AAE65785 Human Sns
63	16	57.1	1956	4	AAE61996 Human vol
64	16	57.1	1956	6	ABG76193 Human per
65	16	57.1	1956	6	ABG75945 Human per
66	16	57.1	1956	6	ABP72253 Human P33
67	16	57.1	1956	6	ADA50152 Human per
68	16	57.1	1956	7	ADJ68903 Human hea
69	16	57.1	1956	8	ADF08471 Human Nav
70	16	57.1	2000	5	ABR806027 Human sod
71	16	57.1	2000	8	ADK81762 Human Nav
72	15	53.6	130	5	ADK35245 Novel hum
73	14	50.0	1366	8	ADJ68488 Human hea
74	14	50.0	1366	8	ADL06575 Human tnm
75	13	46.4	1956	4	AAE65783 Rat Sns1
76	13	46.4	1956	4	AAE61995 Rat perip
77	13	46.4	1956	6	ABG75944 Rat perip
78	13	46.4	1956	6	ADA50144 Rat perip
79	13	46.4	1956	6	ADA50153 Rat perip
80	13	46.4	1957	2	AAE21740 Variant r
81	13	46.4	1957	2	AAE21737 Wild type
82	13	46.4	1957	6	ABG76191 Rat vola
83	13	46.4	1957	6	ADA50156 Rat perip
84	13	46.4	1957	6	ADA50155 Rat perip
85	13	46.4	1957	7	ADA44754 Rat Prote
86	13	46.4	1957	8	ADF08469 Rat Nav 1
87	13	46.4	1958	4	AAE65784 Mouse Sns
88	13	46.4	1962	4	AAE17250 Mouse poly
89	13	46.4	2132	2	AAE21739 Variant r
90	12	42.9	1024	5	ABR80460 LDL recep
91	12	42.9	1233	4	AAE20125 Human sod
92	12	42.9	1243	2	AAE06598 Human sod
93	12	42.9	1243	4	AAE20126 Human sod
94	12	42.9	1444	5	AAO14926 Human sod
95	12	42.9	1791	4	AAE20121 Human sod
96	12	42.9	1791	4	AAO14925 Human sod
97	12	42.9	1791	7	ADD32194 Human Na
98	11	39.3	442	4	ABG04326 Novel hum

CC peptide, or a NHP fusion protein to the body. Nucleotide construct
CC encoding NHP products are also useful in gene therapy for modulating NHP
CC expression and to produce genetically engineered host cells to express
CC NHP products in vivo. NHP nucleotide sequences may also be used as part
CC of ribozyme and/or tripe helix sequences that are useful for NHP gene
CC regulation. The NHP polypeptides are useful for generating antibodies, as
CC reagents in diagnostic assays, for identifying other cellular gene
CC products related to NHP and as reagents in assays for screening for
CC compounds that are useful in the treatment of mental, biological or
CC medical disorders and diseases

XX SQ Sequence 1381 AA;

Query Match 100.0%; Score 28; DB 5; Length 1381;
Best Local Similarity 100.0%; Pred. No. 1.4e-18;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 YMIFPVLVIFLGSFYLINILIAVAMAY 28
Db 399 YMIFPVLVIFLGSFYLINILIAVAMAY 426

RESULT 3
AAE20514
ID AAE20514 standard; protein; 1387 AA.
XX
AC AAE20514;
XX
DT 01-JUL-2002 (first entry)
XX
DE Human ion channel protein #5.
XX
KW Human; novel human protein; NHP; voltage-gated sodium channel;
KW gene therapy; bioreactor; mental disorder; biological disorder;
KW medical disorder.
XX
OS Homo sapiens.
XX

XX Key Location/Qualifiers
FH Misc-difference 981 /note= "Encoded by MTG"
FT 1056
FT Misc-difference /note= "Encoded by RCA"
XX
XX MO200214498-A2.
XX
XX 21-FEB-2002.
XX
XX 15-AUG-2001; 2001WO-US025650.
XX
XX 16-AUG-2000; 2000US-0225989P.
XX
XX (LEXI-) LEXICON GENETICS INC.
XX
XX Turner CA, Mathur B, Mathur D;
XX
XX MPI: 2002-280757/32.
XX
XX N-PSDB; AAD32843.
XX

XX Novel polynucleotides encoding human sodium channel proteins,
XX particularly voltage-gated sodium channel proteins useful for drug
XX screening, diagnosis and in gene therapy of biological disorders.
XX
XX Claim 1; Page 52-55; 83pp; English.

XX The present sequence is novel human protein (NHP), ion channel protein.
XX NHP share structural similarity with mammalian sodium channel proteins
XX particularly voltage-gated sodium channel proteins. NHP oligonucleotides
XX are useful as hybridisation probes for screening libraries and assessing
XX gene expression patterns. Sequences derived from regions adjacent to the
XX intron/exon boundaries of NHP gene can be used to design primers for use
XX in amplification assays to detect mutations within the exons, splice
XX sites, introns that can be used in diagnostics and pharmacogenomics. NHP

CC nucleotide sequences are useful for drug screening effective in the
CC treatment of symptomatic or phenotypic manifestations of perturbing the
CC normal function of NHP in the body, and nucleotide constructs encoding
CC NHP products are useful to genetically engineer host cells to express NHP
CC products in vivo. These genetically engineered cells function as
CC bioreactors in the body delivering a continuous supply of a NHP, a NHP
CC peptide, or a NHP fusion protein to the body. Nucleotide construct
CC encoding NHP products are also useful in gene therapy for modulating NHP
CC expression and to produce genetically engineered host cells to express
CC NHP products in vivo. NHP nucleotide sequences may also be used as part
CC of ribozyme and/or tripe helix sequences that are useful for NHP gene
CC regulation. The NHP polypeptides are useful for generating antibodies, as
CC reagents in diagnostic assays, for identifying other cellular gene
CC products related to NHP and as reagents in assays for screening for
CC compounds that are useful in the treatment of mental, biological or
CC medical disorders and diseases

XX SQ Sequence 1387 AA;

Query Match 100.0%; Score 28; DB 5; Length 1387;
Best Local Similarity 100.0%; Pred. No. 1.4e-18;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 YMIFPVLVIFLGSFYLINILIAVAMAY 28
Db 399 YMIFPVLVIFLGSFYLINILIAVAMAY 426

RESULT 4
AAE20518
ID AAE20518 standard; protein; 1392 AA.
XX
AC AAE20518;
XX
DT 01-JUL-2002 (first entry)
XX
DE Human ion channel protein #9.
XX
KW Human; novel human protein; NHP; voltage-gated sodium channel;
KW gene therapy; bioreactor; mental disorder; biological disorder;
KW medical disorder.
XX
OS Homo sapiens.
XX

XX Key Location/Qualifiers
FH Misc-difference 992 /note= "Encoded by MTG"
FT 1067
FT Misc-difference /note= "Encoded by RCA"
XX
XX MO200214498-A2.
XX
XX 21-FEB-2002.
XX
XX 15-AUG-2001; 2001WO-US025650.
XX
XX 16-AUG-2000; 2000US-0225989P.
XX
XX (LEXI-) LEXICON GENETICS INC.
XX
XX Turner CA, Mathur B, Mathur D;
XX
XX MPI: 2002-280757/32.
XX
XX N-PSDB; AAD32847.
XX

XX Novel polynucleotides encoding human sodium channel proteins,
XX particularly voltage-gated sodium channel proteins useful for drug
XX screening, diagnosis and in gene therapy of biological disorders.
XX
XX Claim 1; Page 75-78; 83pp; English.

XX The present sequence is novel human protein (NHP), ion channel protein.
XX NHP share structural similarity with mammalian sodium channel proteins

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RESULT 5
AAE20519
ID      AAE20519  standard; protein; 1398 AA.
XX

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human ion channel protein #10.

kwx	gene therapy; bioreactor; mental disorder.	voltage-gated sodium channel; biological disorder;
xx	medical disorder.	

Homo sapiens.

Misc-difference 992 Location/Qualifiers

TT	/note=	"Encoded by MTG"
TT	Misc-difference	1067
TT	/note=	"Encoded by RCA"
XX	.	

W0200214498-A2.

21-FEB-2002.

15-AUG-2001; 2001WO-US025650.

16-AUG-2000; 2000US-0225989P.

(LEXI-) LEXICON GENETICS INC.

turner CA, Mathur B, Mathur D;

REF: 2002-280757/32.
N-PSDB; AAD32848

Novel polymers

particular voltage-gated sodium channel proteins, particularly encoding human sodium channel proteins, particularly useful for drug

PT screening, diagnosis and in gene therapy of biological disorders.
XX
PS Claim 1; Page 80-83; 83pp; English.
XX

CC NHP share structural similarity with mammalian sodium channel protein.
CC particularly voltage-gated sodium channel proteins. NHP oligonucleotides
CC are useful as hybridisation probes for screening libraries and assessing
CC gene expression patterns. Sequences derived from regions adjacent to the
CC intron/exon boundaries of NHP gene can be used to design primers for use
CC in amplification assays to detect mutations within the exons, splice
CC sites, introns that can be used in diagnostics and pharmacogenomics. NHP
CC nucleotide sequences are useful for drug screening and perturbations in the
CC treatment of symptomatic or phenotypic manifestations of perturbing the
CC normal function of NHP in the body, and nucleotide constructs encoding the
CC NHP products are useful to genetically engineer host cells to express NHP
CC products *in vivo*. These genetically engineered cells function as
CC bioeffectors in the body delivering a continuous supply of a NHP, a NHP
CC peptide, or a NHP fusion protein to the body. Nucleotide construct
CC encoding NHP products are also useful in gene therapy for modulating NHP
CC expression and to produce genetically engineered host cells to express
CC NHP products *in vivo*. NHP nucleotide sequences may also be used as part
CC of ribozyme and/or triple helix sequences that are useful for NHP gene
CC regulation. The NHP polypeptides are useful for generating antibodies, as
CC reagents in diagnostic assays, for identifying other cellular gene
CC products related to NHP and as reagents in assays for screening for
CC compounds that are useful in the treatment of mental, biological or
CC medical disorders and diseases
XX
80 Sequence 1398 AA,

	Query March	100.0%; Score 28; DB 5;
	Best Local Similarity	100.0%; Pred. NO. 1'ee-18;
	Matches 28; Conservative	0; Mismatches 0; Indels 0; Gaps 0
QY	1 YMFIFLVIFGSPFLINLLIAVVMAY	28
Db	399 YMFIFLVIFGSPFLINLLIAVVMAY	426

AAE20512 standard; protein; 1442 AA

01-JUL-2002 (first entry)

Human ion channel protein #3.

gene therapy; bioreactor; mental disorder; biological disorder.

Homo sapiens.

Key	Location/Qualifiers
Misc 3445	

/note= "Encoded by MTG"

/note= "Encoded by RCA"

WO200214498-A2.

21-FEB-2002.

12-AUG-2001; 2001MO-US025650.

18-AUG-2000; 2000US-0225989P.

Turner CA, Mathur B, Mathur D,

XX WPI; 2002-280757/32.
DR N-PSDB; AAD32841.
XX
XX Novel polynucleotides encoding human sodium channel proteins,
PT particularly voltage-gated sodium channel proteins useful for drug
PT screening, diagnosis and in gene therapy of biological disorders.
XX
PS Claim 1, Page 43-46; 83pp; English.
XX
XX The present sequence is novel human protein (NHP), ion channel protein.
CC NHP share structural similarity with mammalian sodium channel proteins
CC particularly voltage-gated sodium channel proteins. NHP oligonucleotides
CC are useful as hybridisation probes for screening libraries and assessing
CC gene expression patterns. Sequences derived from regions adjacent to the
CC intron/exon boundaries of NHP gene can be used to design primers for use
CC in amplification assays to detect mutations within the exons, splice
CC sites, introns that can be used in diagnostics and pharmacogenomics. NHP
CC nucleotide sequences are useful for drug screening effective in the
CC treatment of symptomatic or phenotypic manifestations of perturbing the
CC normal function of NHP in the body, and nucleotide constructs encoding
CC NHP products are useful to genetically engineer host cells to express NHP
CC products in vivo. These genetically engineered cells function as
CC bioreactors in the body delivering a continuous supply of a NHP, a NHP
CC peptide, or a NHP fusion protein to the body. Nucleotide construct
CC encoding NHP products are also useful in gene therapy for modulating NHP
CC expression and to produce genetically engineered host cells to express
CC NHP products in vivo. NHP nucleotide sequences may also be used as part
CC of ribozyme and/or tripe helix sequences that are useful for NHP gene
CC regulation. The NHP polypeptides are useful for generating antibodies, as
CC reagents in diagnostic assays, for identifying other cellular gene
CC products related to NHP and as reagents in assays for screening for
CC compounds that are useful in the treatment of mental, biological or
CC medical disorders and diseases
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SQ Sequence 1442 AA;
XX
Query Match 100.0%; Score 28; DB 5; Length 1442;
Best Local Similarity 100.0%; Pred. No. 1.5e-18;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 YMIFFVLVIFLGSFYLNILILAVVAMAY 28
Db 399 YMIFFVLVIFLGSFYLNILILAVVAMAY 426
XX
RESULT 7
AAE20517
ID AAE20517 standard; protein; 1453 AA.
XX
XX AAE20517;
XX
XX 01-JUL-2002 (first entry)
XX
XX Human ion channel protein #8.
XX
XX Human; novel human protein; NHP; voltage-gated sodium channel;
KM gene therapy; bioreactor; mental disorder; biological disorder;
KM medical disorder.
XX
XX Homo sapiens.
XX
XX Key Location/Qualifiers
FT Misc-difference 992 /note= "Encoded by MTG"
FT Misc-difference 1067 /note= "Encoded by RCA"
FT
XX
XX WO200214498-A2.
XX
XX 21-FEB-2002.
XX
XX 15-AUG-2001; 2001WO-US025650.
XX
XX

XX 16-AUG-2000; 2000US-0225989P.
XX
XX (LEXI-) LEXICON GENETICS INC.
XX
XX Turner CA, Mathur B, Mathur D;
XX
XX WPI; 2002-280757/32.
DR N-PSDB; AAD32846.
XX
XX Novel polynucleotides encoding human sodium channel proteins,
PT particularly voltage-gated sodium channel proteins useful for drug
PT screening, diagnosis and in gene therapy of biological disorders.
XX
PS Claim 1, Page 70-73; 83pp; English.
XX
XX The present sequence is novel human protein (NHP), ion channel protein.
CC NHP share structural similarity with mammalian sodium channel proteins
CC particularly voltage-gated sodium channel proteins. NHP oligonucleotides
CC are useful as hybridisation probes for screening libraries and assessing
CC gene expression patterns. Sequences derived from regions adjacent to the
CC intron/exon boundaries of NHP gene can be used to design primers for use
CC in amplification assays to detect mutations within the exons, splice
CC sites, introns that can be used in diagnostics and pharmacogenomics. NHP
CC nucleotide sequences are useful for drug screening effective in the
CC treatment of symptomatic or phenotypic manifestations of perturbing the
CC normal function of NHP in the body, and nucleotide constructs encoding
CC NHP products are useful to genetically engineer host cells to express NHP
CC products in vivo. These genetically engineered cells function as
CC bioreactors in the body delivering a continuous supply of a NHP, a NHP
CC peptide, or a NHP fusion protein to the body. Nucleotide construct
CC encoding NHP products are also useful in gene therapy for modulating NHP
CC expression and to produce genetically engineered host cells to express
CC NHP products in vivo. NHP nucleotide sequences may also be used as part
CC of ribozyme and/or tripe helix sequences that are useful for NHP gene
CC regulation. The NHP polypeptides are useful for generating antibodies, as
CC reagents in diagnostic assays, for identifying other cellular gene
CC products related to NHP and as reagents in assays for screening for
CC compounds that are useful in the treatment of mental, biological or
CC medical disorders and diseases
XX
SQ Sequence 1453 AA;
XX
Query Match 100.0%; Score 28; DB 5; Length 1453;
Best Local Similarity 100.0%; Pred. No. 1.5e-18;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 YMIFFVLVIFLGSFYLNILILAVVAMAY 28
Db 399 YMIFFVLVIFLGSFYLNILILAVVAMAY 426
XX
RESULT 8
ADB78596
ID ADB78596 standard; protein; 1795 AA.
XX
XX ADB78596;
XX
XX 04-DEC-2003 (first entry)
XX
XX Human sodium channel subunit mutant SEQ ID NO:140.
XX
XX Human; mutant; ion channel; ion channel subunit; ICS; nootropic;
KM neuroprotective; inotropic; antihypertic; antiarrhythmic; antimigraine;
KM antidepressant; antiparkinsonian; neuroleptic; tranquiliser; analgesic;
KM nephrotoxic; antidiabetic; ophthalmological; epilepsy;
KM ion channel dysfunction; human.
XX
XX Synthetic.
XX Homo sapiens.
XX
XX WO2003008574-A1.
XX
XX

PD 30-JAN-2003.
 XX
 PF 08-JUL-2002; 2002MO-AU000910.
 PR 18-JUL-2001; 2001AU-00006452.
 PR 05-MAR-2002; 2002AU-00000910.
 PR 13-MAY-2002; 2002AU-00002292.
 XX
 PA (BION-) BIONOMICS LTD.
 PA (WALL/) WALLACE R W.
 XX
 PI Mulley JC, Harkin LA, Dibbens LM, Phillips HA, Heron SE;
 PI Berkovic SF, Scheffer IE;
 DR N-PSDB; ADB78635.
 DR WPI: 2003-239332/23.
 XX
 PT Identifying predisposition to an ion channel dysfunction, such as
 PT periodic paralysis, cardiac arrhythmias, migraine, Alzheimer's disease,
 PT schizophrenia, anxiety and depression, by detecting encoding-gene
 PT mutation events.
 XX
 PS Claim 13; SEQ ID NO 140; 106bp; English.
 CC The invention relates to a novel method for identifying a subject
 CC predisposed to a disorder associated with ion channel dysfunction. The
 CC method comprises ascertaining if at least one of the genes encoding ion
 CC channel subunits (ICS) has undergone a mutation event so that a cDNA
 CC derived from the subject has any of 134 nucleotide sequences. The method
 CC of the invention has nootropic, neuroprotective, inotropic, antipyretic,
 CC antirhythmic, antinigraine, antidepressant, antiparkinsonian,
 CC analgesic, tranquiliser, analgesic, nephrotoxic, antidiabetic, and
 CC ion channel agonist, or ion channel antagonist. The methods, isolated
 CC nucleic acids, polypeptides, antibody, selective agonist, antagonist or
 CC modulator of an ion channel, cells and genetically modified non-human
 CC animal, are useful for the diagnosis and treatment of epilepsy and/or a
 CC disorder associated with ion channel dysfunction, such as hyper- or hypo-
 CC kalemic periodic paralysis, myotonias, malignant hyperthermia,
 CC myasthenia, cardiac arrhythmias, episodic ataxia, migraine, Alzheimer's
 CC depression, Parkinson's disease, schizophrenia, hyperphlexia, anxiety,
 CC pain, chronic/acute pain, Bartter's syndrome, polycystic kidney disease,
 CC Dent's disease, hyperinsulinaemic hypoglycaemia of infancy, cystic
 CC fibrosis, congenital stationary night blindness and total colour
 CC blindness. The present sequence represents a mutant protein of the
 CC invention. The sequence data for this patent is not represented in the
 CC printed specification, but was obtained in electronic format directly
 CC from WIPO at ftp.wipo.int/pub/published_pat_sequences.
 XX
 SQ Sequence 1795 AA;
 Query Match 100.0%; Score 28; DB 7; Length 1795;
 Best Local Similarity 100.0%; Pred. No. 1.1e-18;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 YMFVFLVIFLGSFYLNILAVVAMAY 28
 Db 399 YMFVFLVIFLGSFYLNILAVVAMAY 426
 RESULT 9
 ID ADB78597
 XX ADB78597 standard; protein, 1855 AA.
 AC ADB78597;
 XX
 DT 04-DEC-2003 (first entry)
 XX
 DE Human sodium channel subunit mutant SEQ ID NO:141.
 XX
 KW mutein; mutant; ion channel; ion channel subunit; ICS; nootropic;
 KW neuroprotective; inotropic; antipyretic; antirhythmic; antinigraine;

KW antidepressant; antiparkinsonian; neuroleptic; tranquiliser; analgesic;
 KW nephrotoxic; antidiabetic; ophthalmological; epilepsy;
 KW ion channel dysfunction; human.
 XX
 OS Synthetic.
 OS Homo sapiens.
 XX
 PW WO2003008574-A1.
 XX
 PD 30-JAN-2003.
 XX
 PF 08-JUL-2002; 2002MO-AU000910.
 PR 18-JUL-2001; 2001AU-00006452.
 PR 05-MAR-2002; 2002AU-00000910.
 PR 13-MAY-2002; 2002AU-00002292.
 XX
 PA (BION-) BIONOMICS LTD.
 PA (WALL/) WALLACE R W.
 XX
 PI Mulley JC, Harkin LA, Dibbens LM, Phillips HA, Heron SE;
 PI Berkovic SF, Scheffer IE;
 DR N-PSDB; ADB78636.
 DR WPI: 2003-239332/23.
 XX
 PT Identifying predisposition to an ion channel dysfunction, such as
 PT periodic paralysis, cardiac arrhythmias, migraine, Alzheimer's disease,
 PT schizophrenia, anxiety and depression, by detecting encoding-gene
 PT mutation events.
 XX
 PS Claim 13; SEQ ID NO 141; 106bp; English.
 CC The invention relates to a novel method for identifying a subject
 CC predisposed to a disorder associated with ion channel dysfunction. The
 CC method comprises ascertaining if at least one of the genes encoding ion
 CC channel subunits (ICS) has undergone a mutation event so that a cDNA
 CC derived from the subject has any of 134 nucleotide sequences. The method
 CC of the invention has nootropic, neuroprotective, inotropic, antipyretic,
 CC antirhythmic, antinigraine, antidepressant, antiparkinsonian,
 CC analgesic, tranquiliser, analgesic, nephrotoxic, antidiabetic, and
 CC ion channel agonist, or ion channel antagonist. The methods, isolated
 CC nucleic acids, polypeptides, antibody, selective agonist, antagonist or
 CC modulator of an ion channel, cells and genetically modified non-human
 CC animal, are useful for the diagnosis and treatment of epilepsy and/or a
 CC disorder associated with ion channel dysfunction, such as hyper- or hypo-
 CC kalemic periodic paralysis, myotonias, malignant hyperthermia,
 CC myasthenia, cardiac arrhythmias, episodic ataxia, migraine, Alzheimer's
 CC depression, Parkinson's disease, schizophrenia, hyperphlexia, anxiety,
 CC pain, chronic/acute pain, Bartter's syndrome, polycystic kidney disease,
 CC Dent's disease, hyperinsulinaemic hypoglycaemia of infancy, cystic
 CC fibrosis, congenital stationary night blindness and total colour
 CC blindness. The present sequence represents a mutant protein of the
 CC invention. The sequence data for this patent is not represented in the
 CC printed specification, but was obtained in electronic format directly
 CC from WIPO at ftp.wipo.int/pub/published_pat_sequences.
 XX
 SQ Sequence 1855 AA;
 Query Match 100.0%; Score 28; DB 7; Length 1855;
 Best Local Similarity 100.0%; Pred. No. 1.8e-18;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 YMFVFLVIFLGSFYLNILAVVAMAY 28
 Db 399 YMFVFLVIFLGSFYLNILAVVAMAY 426
 RESULT 10
 ID AAE20511
 XX AAE20511 standard; protein, 1962 AA.

XX AAE20511;
AC
XX 01-JUL-2002 (first entry)
DT
XX Human ion channel protein #2.
DE
XX Human; novel human protein; NHP; voltage-gated sodium channel;
KW gene therapy; bioreactor; mental disorder; biological disorder;
KW medical disorder.
XX Homo sapiens.
OS
XX
FH Key Location/Qualifiers
FT Misc-difference 981 /note= "Encoded by MTG"
FT Misc-difference 1056 /note= "Encoded by RCA"
FT
XX MO20021498-AZ.
XX
XX 21-FEB-2002.
XX
XX 15-AUG-2001; 2001WO-US025650.
XX
XX 16-AUG-2000; 2000US-0225989P.
XX
XX (LEXI-) LEXICON GENETICS INC.
XX
XX Turner CA, Mathur B, Mathur D;
XX
XX MPI; 2002-280757/32.
XX
XX N-PSDB; AAD32840.
XX
XX Novel polynucleotides encoding human sodium channel proteins,
PT particularly voltage-gated sodium channel proteins useful for drug
PT screening, diagnosis and in gene therapy of biological disorders.
XX
XX Claim 1; Page 37-41; 83pp; English.

XX The present sequence is novel human protein (NHP), ion channel protein.
CC NHP share structural similarity with mammalian sodium channel proteins
CC particularly voltage-gated sodium channel proteins. NHP oligonucleotides
CC are useful as hybridisation probes for screening libraries and assessing
CC gene expression patterns. Sequences derived from regions adjacent to the
CC intron/exon boundaries of NHP gene can be used to design primers for use
CC in amplification assays to detect mutations within the exons, splice
CC sites, introns that can be used in diagnostics and pharmacogenomics. NHP
CC nucleotide sequences are useful for drug screening effective in the
CC treatment of symptomatic or phenotypic manifestations of perturbing the
CC normal function of NHP in the body, and nucleotide constructs encoding the
CC NHP products are useful to genetically engineer host cells to express NHP
CC products in vivo. These genetically engineered cells function as
CC bioreactors in the body delivering a continuous supply of a NHP, a NHP
CC peptide, or a NHP fusion protein to the body. Nucleotide construct
CC encoding NHP products are also useful in gene therapy for modulating NHP
CC expression and to produce genetically engineered host cells to express
CC NHP products in vivo. NHP nucleotide sequences may also be used as part
CC of ribozyme and/or tripe helix sequences that are useful for NHP gene
CC regulation. The NHP polypeptides are useful for generating antibodies, as
CC reagents in diagnostic assays, for identifying other cellular gene
CC products related to NHP and as reagents in assays for screening for
CC compounds that are useful in the treatment of mental, biological or
CC medical disorders and diseases
XX
XX Sequence 1962 AA;
SQ

Query Match 100.0%; Score 28; DB 5; Length 1962;
Best Local Similarity 100.0%; Pred. No. 1.9e-18;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 YMIFFVLVIFLGSFYLNILAVVAMAY 28
|||||

DB 399 YMIFFVLVIFLGSFYLNILAVVAMAY 426
RESULT 11
AAE20516
ID AAE20516 standard; protein; 1973 AA.
XX
XX AAE20516;
AC
XX 01-JUL-2002 (first entry)
DT
XX Human ion channel protein #7.
DE
XX Human; novel human protein; NHP; voltage-gated sodium channel;
KW gene therapy; bioreactor; mental disorder; biological disorder;
KW medical disorder.
XX Homo sapiens.
OS
XX
FH Key Location/Qualifiers
FT Misc-difference 992 /note= "Encoded by MTG"
FT Misc-difference 1067 /note= "Encoded by RCA"
FT
XX MO20021498-AZ.
XX
XX 21-FEB-2002.
XX
XX 15-AUG-2001; 2001WO-US025650.
XX
XX 16-AUG-2000; 2000US-0225989P.
XX
XX (LEXI-) LEXICON GENETICS INC.
XX
XX Turner CA, Mathur B, Mathur D;
XX
XX MPI; 2002-280757/32.
XX
XX N-PSDB; AAD32845.
XX
XX Novel polynucleotides encoding human sodium channel proteins,
PT particularly voltage-gated sodium channel proteins useful for drug
PT screening, diagnosis and in gene therapy of biological disorders.
XX
XX Claim 1; Page 64-68; 83pp; English.

XX The present sequence is novel human protein (NHP), ion channel protein.
CC NHP share structural similarity with mammalian sodium channel proteins
CC particularly voltage-gated sodium channel proteins. NHP oligonucleotides
CC are useful as hybridisation probes for screening libraries and assessing
CC gene expression patterns. Sequences derived from regions adjacent to the
CC intron/exon boundaries of NHP gene can be used to design primers for use
CC in amplification assays to detect mutations within the exons, splice
CC sites, introns that can be used in diagnostics and pharmacogenomics. NHP
CC nucleotide sequences are useful for drug screening effective in the
CC treatment of symptomatic or phenotypic manifestations of perturbing the
CC normal function of NHP in the body, and nucleotide constructs encoding the
CC NHP products are useful to genetically engineer host cells to express NHP
CC products in vivo. These genetically engineered cells function as
CC bioreactors in the body delivering a continuous supply of a NHP, a NHP
CC peptide, or a NHP fusion protein to the body. Nucleotide construct
CC encoding NHP products are also useful in gene therapy for modulating NHP
CC expression and to produce genetically engineered host cells to express
CC NHP products in vivo. NHP nucleotide sequences may also be used as part
CC of ribozyme and/or tripe helix sequences that are useful for NHP gene
CC regulation. The NHP polypeptides are useful for generating antibodies, as
CC reagents in diagnostic assays, for identifying other cellular gene
CC products related to NHP and as reagents in assays for screening for
CC compounds that are useful in the treatment of mental, biological or
CC medical disorders and diseases
XX
XX Sequence 1973 AA;
SQ

```

Query Match          100.0%; Score 28; DB 5; Length 1973;
Best Local Similarity 100.0%; Pred. No. 1.9e-18;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 YMIFFVLVIFLGSPYLINLIILAVVAMAY 28
DB 399 YMIFFVLVIFLGSPYLINLIILAVVAMAY 426

RESULT 12
ABR83185
ID ABR83185 standard; protein; 1981 AA.
AC ABR83185;
XX
XX
XX 15-JAN-2004 (first entry)
DE Human SCN1A splice variant -84P:SCN1ADLP654-681.
XX
XX SCN1A; sodium channel type 1 alpha-subunit; anticonvulsant; analgesic;
KM neuroprotective; anesthetic; cytosolic; cerebroprotective; cardiant;
KM hypotensive; gene therapy; human; splice variant.
XX
OS Homo sapiens.
XX
XX WO2003072751-A2.
XX
XX 04-SEP-2003.
XX
XX 25-FEB-2003; 2003WO-US006010.
XX
XX 25-FEB-2002; 2002US-0359382P.
XX
XX (UYVA-) UNIV VANDERBILT.
XX
XX George AL, Losein C;
XX
XX WPI, 2003-712725/67.
XX
XX DR N-PSDB; ACF57880.
XX
XX
XX Recombinantly expressed sodium channel type 1 alpha subunit, useful in
XX screening for modulators, for treating e.g. epilepsy.
XX
XX Disclosure; Page 162-169; 176pp; English.
XX
XX The invention relates to a recombinantly expressed and isolated human
XX SCN1A (sodium channel type 1 alpha-subunit) (I). (I) optionally
XX incorporated into a cell, is used to screen for specific modulators,
XX potentially useful as anticonvulsant, antiepileptic, neuroprotective,
XX analgesic and/or anesthetic agents, e.g. for treating severe myoclonic
XX epilepsy of infancy, stroke, cardiac arrest, hyperkalemic paralysis,
XX motor endplate diseases, hypertension, congestive heart failure and
XX CC muscular dystrophy also to treat cancer (SCN1A is expressed in prostatic
XX CC by gene therapy vectors that express (I) or the modulators. The
XX CC modulators, also antibodies directed against (I), are used to detect
XX CC sodium channel polypeptides. The present sequence represents a human
XX CC SCN1A splice variant 84P:SCN1ADLP654-681
XX
XX Sequence 1981 AA;
XX
XX Query Match          100.0%; Score 28; DB 7; Length 1981;
XX Best Local Similarity 100.0%; Pred. No. 1.9e-18;
XX Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX OY 1 YMIFFVLVIFLGSPYLINLIILAVVAMAY 28
XX DB 399 YMIFFVLVIFLGSPYLINLIILAVVAMAY 426
XX
XX RESULT 13
XX AAE20510
XX ID AAE20510 standard; protein; 1998 AA.

```

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XX
XX AC AAE20510;
XX
XX DT 01-JUL-2002 (first entry)
XX
XX DE Human ion channel protein #1.
XX
XX Human; novel human protein; NHP; voltage-gated sodium channel;
XX KM gene therapy; bioreactor; mental disorder; biological disorder;
XX KM medical disorder.
XX
XX OS Homo sapiens.
XX
XX Key Location/Qualifiers
XX FH MISC-difference 981
XX FT MISC-difference 1056
XX FT MISC-difference 1056
XX FT MISC-difference 1056
XX
XX WO200214498-A2.
XX
XX 21-FEB-2002.
XX
XX 15-AUG-2001; 2001WO-US025650.
XX
XX 16-AUG-2000; 2000US-0225989P.
XX
XX (LEXI-) LEXICON GENETICS INC.
XX
XX Turner CA, Mathur B, Mathur D;
XX
XX WPI; 2002-280757/32.
XX
XX DR N-PSDB; AAD32839.
XX
XX Novel polynucleotides encoding human sodium channel proteins,
XX particularly voltage-gated sodium channel proteins useful for drug
XX screening, diagnosis and in gene therapy of biological disorders.
XX
XX Claim 1; Page 30-34; 83pp; English.
XX
XX The present sequence is novel human protein (NHP), ion channel protein.
XX NHP share structural similarity with mammalian sodium channel proteins.
XX particularly voltage-gated sodium channel proteins. NHP oligonucleotides
XX are useful as hybridisation probes for screening libraries and assessing
XX gene expression patterns. Sequences derived from regions adjacent to the
XX intron/exon boundaries of NHP gene can be used to design primers for use
XX in amplification assays to detect mutations within the exons, splice
XX sites, introns that can be used in diagnostics and pharmacogenomics. NHP
XX nucleotide sequences are useful for drug screening effective in the
XX treatment of symptomatic or phenotypic manifestations of perturbing the
XX normal function of NHP in the body, and nucleotide constructs encoding the
XX NHP products are useful to genetically engineered host cells to express NHP
XX products in vivo. These genetically engineered cells function as
XX bioreactors in the body delivering a continuous supply of a NHP, a NHP
XX peptide, or a NHP fusion protein to the body. Nucleotide construct
XX encoding NHP products are also useful in gene therapy for modulating NHP
XX expression and to produce genetically engineered host cells to express
XX NHP products in vivo. NHP nucleotide sequences may also be used as part
XX of ribozyme and/or triple helix sequences that are useful for NHP gene
XX regulation. The NHP polypeptides are useful for generating antibodies, as
XX reagents in diagnostic assays, for identifying other cellular gene
XX products related to NHP and as reagents in assays for screening for
XX compounds that are useful in the treatment of mental, biological or
XX medical disorders and diseases
XX
XX Sequence 1998 AA;
XX
XX Query Match          100.0%; Score 28; DB 5; Length 1998;
XX Best Local Similarity 100.0%; Pred. No. 1.9e-18;
XX Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX OY 1 YMIFFVLVIFLGSPYLINLIILAVVAMAY 28
XX

```

Db 399 YMIFFVLVIFLGSFYILNLIILAVVAMAY 426

RESULT 14

ABR83184 standard; protein; 1998 AA.

AC ABR83184;

DT 15-JAN-2004 (first entry)

DE Human SCN1A splice variant -33P:SCN1ADELP671-681.

XX SCN1A; sodium channel type 1 alpha-subunit; anticonvulsant; analgesic;

KW neuroprotective; anesthetic; cyostatic; cerebroprotective; cardant;

KM hypotensive; gene therapy; human; splice variant.

XX Homo sapiens.

OS

PN WO2003072751-A2.

PD 04-SEP-2003.

XX

PF 25-FEB-2003; 2003WO-US006010.

XX

PR 25-FEB-2002; 2002US-0359382P.

XX

PA (UYVA-) UNIV VANDERBILT.

XX

PI George AL, Lossin C;

XX

DR MPI: 2003-712725/67.

DR N-PSDB; ACF57879.

XX

PT Recombinantly expressed sodium channel type 1 alpha subunit, useful in

PT screening for modulators, for treating e.g. epilepsy.

XX

PS Disclosure; Page 148-156; 176pp; English.

XX

CC The invention relates to a recombinantly expressed and isolated human

CC SCN1A (sodium channel type 1 alpha-subunit) (I). (I), optionally

CC incorporated into a cell, is used to screen for specific modulators,

CC potentially useful as anticonvulsant, antiepileptic, neuroprotective,

CC analgesic and/or anesthetic agents, e.g. for treating severe myoclonic

CC epilepsy of infancy, stroke, cardiac arrest, hyperkalemic paralysis,

CC motor endplate diseases, hypertension, congestive heart failure and

CC muscular dystrophy also to treat cancer (SCN1A is expressed in prostatic

CC and metastatic cancer cell lines). These activities can also be provided

CC by gene therapy vectors that express (I) or the modulators. The

CC modulators, also antibodies directed against (I), are used to detect

CC sodium channel polypeptides. The present sequence represents a human

CC SCN1A splice variant 33P:SCN1ADELP671-681 encoding cDNA

XX

SQ Sequence 1998 AA;

Query Match 100.0%; Score 28; DB 7; Length 1998;

Best Local Similarity 100.0%; Pred. No. 1.9e-18;

Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 YMIFFVLVIFLGSFYILNLIILAVVAMAY 28

DB 399 YMIFFVLVIFLGSFYILNLIILAVVAMAY 426

RESULT 15

ABR06026 standard; protein; 1999 AA.

AC ABR06026;

DT 10-MAY-2002 (first entry)

XX

DE Human sodium channel SCN1A protein SEQ ID NO:2.

XX

KW Human; sodium channel; SCN1A; chromosome 2q24;

XX familial hypercalcaemic periodic paralysis; motor endplate disease.

XX

OS Homo sapiens.

XX

PN WO200196552-A1.

XX

PD 20-DEC-2001.

XX

PF 12-JUN-2001; 2001WO-JP004956.

XX

PR 13-JUN-2000; 2000JP-00177540.

XX

PR 13-JUN-2000; 2000JP-00177544.

XX

PA (NISC-) JAPAN SCI & TECHNOLOGY CORP.

XX

PI Kanazawa I, Goto J, Jeong S;

XX

DR MPI: 2002-098066/13.

DR N-PSDB; ABL39689.

XX

PS Claim 1; Page 40-49; 88pp; Japanese.

XX

CC The present invention describes human sodium channels SCN1A and SCN3A.

CC CC SCN3A have been located to the human chromosome 2 long arm, positions

CC 2q24 and 2q24-31 respectively. The sodium channel proteins are useful in

CC studying the physiological mechanism in which excitant cells participate

CC and cause human diseases, and in developing remedies for e.g. familial

CC hypercalcaemic periodic paralysis of extremities and motor endplate

CC disease

XX

SQ Sequence 1999 AA;

Query Match 100.0%; Score 28; DB 5; Length 1999;

Best Local Similarity 100.0%; Pred. No. 1.9e-18;

Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 YMIFFVLVIFLGSFYILNLIILAVVAMAY 28

DB 399 YMIFFVLVIFLGSFYILNLIILAVVAMAY 426

RESULT 16

AAB99676 standard; protein; 2005 AA.

AC AAB99676;

DT 04-SEP-2001 (first entry)

XX

DE Human adult form of SCN2A protein sequence SEQ ID NO:35.

XX

KW Human; epilepsy; chromosome 2; SCN1A; SCN2A; SCN3A; identification;

KW diagnosis; mutation; chromosome 2q23-q31; neurological disorder;

KM anticonvulsant; neuroprotective.

XX

OS Homo sapiens.

XX

PN WO200138564-A2.

XX

PD 31-MAY-2001.

XX

PF 24-NOV-2000; 2000WO-CA001404.

XX

PR 26-NOV-1999; 99US-0167623P.

XX

PA (UYWC-) UNIV MCGILL.

```

XX Rouleau GA, Lafreniere RG, Rochefort D, Cossette P, Ragedale D;
XX WPI, 2001-355945/37.
XX N-PSDB; AAH55793.
XX
XX Determining a predisposition to epilepsy and/or development of epilepsy
XX comprises determining the genotype of SCN1A, SCN2A and/or SCN3A, or a DNA
XX variant, equivalent, or mutation which shows a linkage disequilibrium.
XX Disclosure; Page 123-130; 268pp; English.
XX
XX The present invention describes a method (M1) of determining an
XX individual's predisposition to epilepsy and/or development of epilepsy,
XX as well as predicting the individual's response to medication. The method
XX comprises determining the genotype of at least one gene selected from
XX SCN1A, SCN2A or SCN3A, or a DNA variant, equivalent, or mutation which
XX shows a linkage disequilibrium. SCN1A, SCN2A and SCN3A are all sodium
XX channel genes located on chromosome 2. The idiopathic generalised
XX epilepsy (IGE) gene is more specifically localised on chromosome 2q23-
XX q31. Compounds identified as modulators of the biological activity of
XX SCN1A, SCN2A or SCN3A proteins or genes, are useful for treating epilepsy
XX or other neurological disorders. They have anticonvulsant and
XX neuroprotective activities. AAH55763 to AAH56164 and AAB99674 to AAB99679
XX represent SCN1A, SCN2A, and SCN3A cDNAs, gene fragments, PCR primers,
XX oligonucleotides and proteins given in the exemplification of the present
XX invention.
XX
SQ Sequence 2005 AA;

Query Match
Best Local Similarity 100.0%; Score 28; DB 4; Length 2005;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 YMIFVLVIFLGSFYILNILLAVVMAY 28
DB 401 YMIFVLVIFLGSFYILNILLAVVMAY 428

RESULT 17
AAB99677
ID AAB99677 standard; protein; 2005 AA.
AC AAB99677;
XX
XX 04-SEP-2001 (first entry)
XX
XX Human neonatal form of SCN2A protein sequence SEQ ID NO:36.
XX
XX Human; epilepsy; chromosome 2; SCN1A; SCN2A; SCN3A; identification;
XX diagnosis; mutation; chromosome 2q23-q31; neurological disorder;
XX anticonvulsant; neuroprotective.
XX
XX Homo sapiens.
XX
XX WO200138564-A2.
XX
XX 31-MAY-2001.
XX
XX 24-NOV-2000; 2000WO-CA001404.
XX
XX 26-NOV-1999; 99US-0167623P.
XX
XX (UMC-) UNITV MCGILL.
XX
XX Rouleau GA, Lafreniere RG, Rochefort D, Cossette P, Ragedale D;
XX WPI, 2001-355945/37.
XX N-PSDB; AAH55794.
XX
XX Determining a predisposition to epilepsy and/or development of epilepsy
XX comprises determining the genotype of SCN1A, SCN2A and/or SCN3A, or a DNA
XX variant, equivalent, or mutation which shows a linkage disequilibrium.
XX
XX

```

```

XX Disclosure; Page 131-138; 268pp; English.
XX
XX The present invention describes a method (M1) of determining an
XX individual's predisposition to epilepsy and/or development of epilepsy,
XX as well as predicting the individual's response to medication. The method
XX comprises determining the genotype of at least one gene selected from
XX SCN1A, SCN2A or SCN3A, or a DNA variant, equivalent, or mutation which
XX shows a linkage disequilibrium. SCN1A, SCN2A and SCN3A are all sodium
XX channel genes located on chromosome 2. The idiopathic generalised
XX epilepsy (IGE) gene is more specifically localised on chromosome 2q23-
XX q31. Compounds identified as modulators of the biological activity of
XX SCN1A, SCN2A or SCN3A proteins or genes, are useful for treating epilepsy
XX or other neurological disorders. They have anticonvulsant and
XX neuroprotective activities. AAH55763 to AAH56164 and AAB99674 to AAB99679
XX represent SCN1A, SCN2A, and SCN3A cDNAs, gene fragments, PCR primers,
XX oligonucleotides and proteins given in the exemplification of the present
XX invention.
XX
SQ Sequence 2005 AA;

Query Match
Best Local Similarity 100.0%; Score 28; DB 4; Length 2005;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 YMIFVLVIFLGSFYILNILLAVVMAY 28
DB 401 YMIFVLVIFLGSFYILNILLAVVMAY 428

RESULT 18
AAB83627
ID AAB83627 standard; protein; 2005 AA.
AC AAB83627;
XX
XX 10-OCT-2002 (first entry)
XX
XX Human GEFs+ protein with SCN2A mutation.
XX
XX Human; GEFs+; SCN2A; mutant; mutein;
XX generalized epilepsy with febrile seizure plus.
XX
XX Homo sapiens.
XX
XX JP2002136289-A.
XX
XX 14-MAY-2002.
XX
XX 01-NOV-2000; 2000JP-0034969.
XX
XX 01-NOV-2000; 2000JP-0034969.
XX
XX 01-NOV-2000; 2000JP-0034969.
XX
XX (KAGA-) KAGAKU GIJUTSU SHINKO JIGYODAN.
XX (RIKA) RIKAGAKU KENKYUSHO.
XX
XX WPI: 2002-552308/59.
XX N-PSDB; ABO79201.
XX
XX A human polynucleotide which is complementary to an mRNA transcribed from
XX useful for diagnosing GEFs+.
XX
XX Claim 10; Page 29-34; 37pp; Japanese.
XX
XX This invention relates to a human polynucleotide which is complementary
XX to an mRNA transcribed from a "generalized epilepsy with febrile seizure
XX plus" (GEFS+)-related gene. The gene is useful for diagnosing GEFs+. The
XX present sequence represents the human GEFs+ protein sequence with SCN2A
XX mutation
XX
XX Sequence 2005 AA;

```

Query Match 100.0%; Score 28; DB 5; Length 2005;
 Best Local Similarity 100.0%; Pred. No. 1.9e-18;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 YMIFFVLVIFLGSFYILNIIILAVVMAY 28
 |||||
 DB 401 YMIFFVLVIFLGSFYILNIIILAVVMAY 428

RESULT 19

ADB78604
 ID ADB78604 standard; protein; 2005 AA.

XX ADB78604;
 AC
 XX

DT 04-DEC-2003 (first entry)

DE Human sodium channel subunit mutant SEQ ID NO:148.

XX muretin; mutant; ion channel; ion channel subunit; ICS; noctropic;
 XX neuroprotective; inotropic; antipyrctic; antiarrhythmic; antimigraine;
 KM antidepressant; antiparkinsonian; neuroleptic; tranquiliser; analgesic;
 KM nephrotropic; antidiabetic; ophthalmological; epilepsy;
 KM ion channel dysfunction; human.

OS Synthetic.
 OS Homo sapiens.

XX WO2003008574-A1.

XX 30-JAN-2003.

XX 08-JUL-2002; 2002WO-AU000910.

XX 18-JUL-2001; 2001AU-00006452.

XX 05-MAR-2002; 2002AU-00000910.

XX 13-MAY-2002; 2002AU-00002292.

XX (BION-) BIONOMICS LTD.
 PA (WALL) WALLACE R W.

XX Mulley JC, Harkin LA, Dibbens LM, Phillips HA, Heron SE;
 PI Berkovic SF, Scheffer IE;

XX WPI; 2003-239332/23.
 DR N-PSDB; ADB78643.

PT Identifying predisposition to an ion channel dysfunction, such as
 PT periodic paralysis, cardiac arrhythmias, migraine, Alzheimer's disease,
 PT schizophrenia, anxiety and depression, by detecting encoding-gene
 PT mutation events.

PS Claim 13; SEQ ID NO 148; 106pp; English.

CC The invention relates to a novel method for identifying a subject
 CC predisposed to a disorder associated with ion channel dysfunction. The
 CC method comprises ascertaining if at least one of the genes encoding ion
 CC channel subunits (ICS) has undergone a mutation event so that a cDNA
 CC derived from the subject has any of 134 nucleotide sequences. The method
 CC of the invention has noctropic, neuroprotective, inotropic, antipyrctic,
 CC antiarrhythmic, antiemetic, antidepressant, antiparkinsonian,
 CC neuroleptic, tranquiliser, analgesic, nephrotropic, antidiabetic, and
 CC ophthalmological activity. A polynucleotide of the invention acts as an
 CC ion channel agonist, or ion channel antagonist. The methods, isolated
 CC nucleic acids, polypeptides, antibody, selective agonist, antagonist or
 CC modulator of an ion channel, cells and genetically modified non-human
 CC animal, are useful for the diagnosis and treatment of epilepsy and/or a
 CC disorder associated with ion channel dysfunction, such as hyper- or hypo-
 CC kalemic periodic paralysis, myotonia, malignant hyperthermia,
 CC myasthenia, cardiac arrhythmias, episodic ataxia, migraine, Alzheimer's
 CC disease, Parkinson's disease, schizophrenia, hyperplexia, anxiety,
 CC depression, phobic obsessive symptoms, neuropathic pain, inflammatory
 CC pain, chronic/acute pain, Bartter's syndrome, polycystic kidney disease,

CC Dent's disease, hyperinsulinaemic hypoglycaemia of infancy, cystic
 CC fibrosis, congenital stationary night blindness and total colour
 CC blindness. The present sequence represents a mutant protein of the
 CC invention. The sequence data for this patent is not represented in the
 CC printed specification, but was obtained in electronic format directly
 CC from WIPO at ftp.wipo.int/pub/published_pat_sequences.

XX SQ Sequence 2005 AA;

Query Match 100.0%; Score 28; DB 7; Length 2005;
 Best Local Similarity 100.0%; Pred. No. 1.9e-18;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 YMIFFVLVIFLGSFYILNIIILAVVMAY 28
 |||||
 DB 401 YMIFFVLVIFLGSFYILNIIILAVVMAY 428

RESULT 20

ADB78603
 ID ADB78603 standard; protein; 2005 AA.

XX ADB78603;
 AC
 XX

DT 04-DEC-2003 (first entry)

DE Human sodium channel subunit mutant SEQ ID NO:147.

XX muretin; mutant; ion channel; ion channel subunit; ICS; noctropic;
 XX neuroprotective; inotropic; antipyrctic; antiarrhythmic; antimigraine;
 KM antidepressant; antiparkinsonian; neuroleptic; tranquiliser; analgesic;
 KM nephrotropic; antidiabetic; ophthalmological; epilepsy;
 KM ion channel dysfunction; human.

OS Synthetic.
 OS Homo sapiens.

XX WO2003008574-A1.

XX 30-JAN-2003.

XX 08-JUL-2002; 2002WO-AU000910.

XX 18-JUL-2001; 2001AU-00006452.

XX 05-MAR-2002; 2002AU-00000910.

XX 13-MAY-2002; 2002AU-00002292.

XX (BION-) BIONOMICS LTD.
 PA (WALL) WALLACE R W.

XX Mulley JC, Harkin LA, Dibbens LM, Phillips HA, Heron SE;
 PI Berkovic SF, Scheffer IE;

XX WPI; 2003-239332/23.
 DR N-PSDB; ADB78642.

PT Identifying predisposition to an ion channel dysfunction, such as
 PT periodic paralysis, cardiac arrhythmias, migraine, Alzheimer's disease,
 PT schizophrenia, anxiety and depression, by detecting encoding-gene
 PT mutation events.

PS Claim 13; SEQ ID NO 147; 106pp; English.

CC The invention relates to a novel method for identifying a subject
 CC predisposed to a disorder associated with ion channel dysfunction. The
 CC method comprises ascertaining if at least one of the genes encoding ion
 CC channel subunits (ICS) has undergone a mutation event so that a cDNA
 CC derived from the subject has any of 134 nucleotide sequences. The method
 CC of the invention has noctropic, neuroprotective, inotropic, antipyrctic,
 CC antiarrhythmic, antiemetic, antidepressant, antiparkinsonian,
 CC neuroleptic, tranquiliser, analgesic, nephrotropic, antidiabetic, and
 CC ophthalmological activity. A polynucleotide of the invention acts as an
 CC ion channel agonist, or ion channel antagonist. The methods, isolated

	Query Match	100.0%;	Score 28;	DB 7;	Length 2005;
	Best Local Similarity	100.0%;	Pred.	No. 1.9e-18;	
	Matches	28;	Conservative	0;	Mismatches 0;
					Indels 0; Gaps 0;
Qy					
	1 YMFIFVLVIFFGSEFLINLLIAYVAMAY				28
Dd	401 YMFIIVLVIFFGSEFLINLLIAYVAMAY				428

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RESULT 21
ADB78605
ID      ADB78605 standard; protein; 2005 AA.
xy

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DT 04-DEC-2003 (first entry)
XX

mutant subunit channel

neurotransmitter; ion channel; ion channel subunit; ICS; nootropic; neuroprotective; inotropic; antipyretic; antirhythmic; antimigraine; antidepressant; antiparkinsonian; neuroleptic; tranquiliser; analgesic; nephrotoxic; antidiabetic; ophthalmological; epilepsy; ion channel dysfunction; human.

Synthetic.
Homo sapiens.

WO2003008574-A1

30-JAN-2003.

08-JUL-2002; 2002WO-AU000910.

18-JUL-2001; 2001AU-00006452.

13-MAY-2002; 2002AU-00002292.

A (BLON-) BIONOMICS LTD.
A (WALF.) WAITAGE & SONS LTD.

Muller, J.C. Harkins

BEKROVIC SR, Schetter IE;

N-PSDB; ADB78644

Identifying predisposing

schizophrenia, anxiety

X

claim 13; SEQ ID NO 149; 106pp; English.

... invention relates to a novel method for identifying a subject

CC predisposed to a disorder associated with ion channel dysfunction. The
CC method comprises ascertaining if at least one of the genes encoding ion
CC channel subunits (ICS) has undergone a mutation event so that a cDNA
CC derived from the subject has any of 13 nucleotide sequences. The method
CC of the invention has nucleotide, neuroprotective, inotropic, antihypertic,
CC antiarhythmic, antimigraine, antidepressant, antiparkinsonian,
CC neuroleptic, triaquilliser, analgesic, nephrotoxic, antidiabetic, and
CC ophthalmological activity. A polynucleotide of the invention acts as an
CC ion channel agonist, or ion channel antagonist. The methods, isolated
CC nucleic acids, polypeptides, antibody, sequesterist. The methods, isolated
CC modulator of an ion channel, cells and genetically modified non-human
CC animal, are useful for the diagnosis and treatment of epilepsy and/or a
CC disorder associated with ion channel dysfunction, such as hyper- or hypo-
CC kalemic periodic paralysis, myotonia, malignant hyperthermia,
CC myaesthesia, cardiac arrhythmias, episodic ataxia, migraine, Alzheimer's
CC disease, Parkinson's disease, schizophrenia, hyperkplexia, anxiety,
CC depression, phobic obsessive symptoms, neuropathic pain, inflammatory
CC pain, chronic/acute pain, Bartter's syndrome, poly cystic kidney disease,
CC Dent's disease/hyponatraemic hypoglycaemia of infancy, cystic
CC fibrosis, congenital stationary night blindness and total colour
CC blindness. The present sequence represents a mutant protein of the
CC invention. The sequence data for this patent is not represented in the
CC printed specification, but was obtained in electronic format directly
CC from WIPO at ftp.wipo.int/pub/published_pat_sequences.
XX

	Query Match	100.0%	Score 28 ; DB 7;	Length 2005;
	Best Local Similarity	100.0% ; Pred.	No. 1_9e-18;	
	Matches	28 ; Conservative	0 ; Mismatches	0 ; Indels
	Gaps	0 ; Gaps		
Oy	I YNIFPVLVFLGSPYLNMLIAVAMAY	28		
Dd	I NIFFPVLVFGLSPLYNLILAVAMAY	428		

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RESULT 22
ADC46947
ID      ADC46947 standard; protein; 2005 AA
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AC	ADC46947
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DI 18-DEC-2003 (first entry)
XX

DE XX	human SCNA amino acid sequence #SEQ ID 3
1	1
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100	100

SCN2A; voltage-gated ion channel; human; neuroprotective; gene therapy, vaccine: Alzheimer's disease

Homo sapiens

PN WO2003060525-A1

PD 24-JUL-2003

PF 16-JAN-2003; 2003WO-EP0000400

17-JAN-2002; 2002EP~00001236

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TOTAL			

N-PSDB; ADC46961.

Diagnosing or prognosticating a neurodegenerative disease by detecting

gene coding for the voltage-gated ion channel SCN2A

Disclosure; Fig 9; 67pp; English.

CC The invention relates to a method for diagnosing or prognosticating a
 CC neurodegenerative disease in a subject, or determining whether a subject
 CC is at increased risk of developing the disease. The method comprises
 CC detecting the level and/or activity of a transcription or translation
 CC product of the gene coding for the voltage-gated ion channel SCN2A. The
 CC modulator of an activity and/or of a level of at least one substrate is
 CC useful for preparing a composition for treating or preventing a
 CC neurodegenerative disease, in particular Alzheimer's disease. The current
 CC sequence represents the human SCN2A amino acid sequence.

XX
 SQ Sequence 2005 AA;

Query Match 100.0%; Score 28; DB 7; Length 2005;
 Best Local Similarity 100.0%; Pred. No. 1.9e-18;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 YMIFVLVIFLGSFYLINILAVVMAY 28
 |||||
 DB 401 YMIFVLVIFLGSFYLINILAVVMAY 428

RESULT 23

AAB9674 ID AAB9674 standard; protein; 2009 AA.

XX AAB9674;

DT 04-SEP-2001 (first entry)

DE Human adult form of SCN1A protein sequence SEQ ID NO:3.

XX Human; epilepsy; chromosome 2; SCN1A; SCN2A; SCN3A; identification;
 KW diagnosis; mutation; chromosome 2q23-q31; neurological disorder;
 KW anticonvulsant; neuroprotective.

XX Homo sapiens.

OS WO200138564-A2.

PN 31-MAY-2001.

XX 24-NOV-2000; 2000WO-CA001404.

XX 26-NOV-1999; 99US-0167623P.

XX (UWMC-) UNIV MCGILL.

PI Rouleau GA, Lafreniere RG, Rochefort D, Cossette P, Ragsdale D;

DR WPI; 2001-355945/37.

XX N-PSDB; AAH55763.

PT Determining a predisposition to epilepsy and/or development of epilepsy
 PT comprises determining the genotype of SCN1A, SCN2A and/or SCN3A, or a DNA
 PT variant, equivalent, or mutation which shows a linkage disequilibrium.

XX Disclosure: Page 96-104; 268pp; English.

XX The present invention describes a method (M1) of determining an
 CC individual's predisposition to epilepsy and/or development of epilepsy,
 CC as well as predicting the individual's response to medication. The method
 CC comprises determining the genotype of at least one gene selected from
 CC SCN1A, SCN2A or SCN3A, or a DNA variant, equivalent, or mutation which
 CC shows a linkage disequilibrium. SCN1A, SCN2A and SCN3A are all sodium
 CC channel genes located on chromosome 2. The idiopathic generalised
 CC epilepsy (IGE) gene is more specifically localised on chromosome 2q23-
 CC q31. Compounds identified as modulators of the biological activity of
 CC SCN1A, SCN2A or SCN3A proteins or genes, are useful for treating epilepsy
 CC or other neurological disorders. They have anticonvulsant and
 CC neuroprotective activities. AAH55763 to AAH56164 and AAB9674 to AAB9679
 CC represent SCN1A, SCN2A, and SCN3A cDNAs, gene fragments, PCR primers,
 CC oligonucleotides and proteins given in the exemplification of the present
 CC invention

XX
 SQ Sequence 2009 AA;

Query Match 100.0%; Score 28; DB 4; Length 2009;
 Best Local Similarity 100.0%; Pred. No. 1.9e-18;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 YMIFVLVIFLGSFYLINILAVVMAY 28
 |||||
 DB 399 YMIFVLVIFLGSFYLINILAVVMAY 426

RESULT 24
 AAE20515

ID AAE20515 standard; protein; 2009 AA.

XX AAE20515;

DT 01-JUL-2002 (first entry)

DE Human ion channel protein #6.

XX Human; novel human protein; NHP; voltage-gated sodium channel;
 KW gene therapy; bioreactor; mental disorder; biological disorder;
 KW medical disorder.

XX Homo sapiens.

FN Key Location/Qualifiers

FT Misc-difference 992 /note= "Encoded by MTG"

FT Misc-difference 1067 /note= "Encoded by RCA"

XX WO200214498-A2.

XX 21-FEB-2002.

XX 15-AUG-2001; 2001WO-US025650.

XX 16-AUG-2000; 2000US-0225989P.

XX (LEXI-) LEXICON GENETICS INC.

PI Turner CA, Mathur B, Mathur D;

DR WPI; 2002-280757/32.

XX N-PSDB; AAD32844.

PT Novel polynucleotides encoding human sodium channel proteins,
 PT particularly voltage-gated sodium channel proteins useful for drug
 PT screening, diagnosis and in gene therapy of biological disorders.

XX Claim 5; Page 57-62; 83pp; English.

XX The present sequence is novel human protein (NHP), ion channel protein.
 CC NHP share structural similarity with mammalian sodium channel proteins.
 CC particularly voltage-gated sodium channel proteins. NHP oligonucleotides
 CC are useful as hybridisation probes for screening libraries and assessing
 CC gene expression patterns. Sequences derived from regions adjacent to the
 CC intron/exon boundaries of NHP gene can be used to design primers for use
 CC in amplification assays to detect mutations within the exons, splice
 CC sites, introns that can be used in diagnostics and pharmacogenomics. NHP
 CC nucleotide sequences are useful for drug screening effective in the
 CC treatment of symptomatic or phenotypic manifestations of perturbing the
 CC normal function of NHP in the body, and nucleotide constructs encoding
 CC NHP products are useful to genetically engineer host cells to express NHP
 CC products in vivo. These genetically engineered cells function as
 CC bioreactors in the body delivering a continuous supply of a NHP, a NHP
 CC peptide, or a NHP fusion protein to the body. Nucleotide construct
 CC encoding NHP products are also useful in gene therapy for modulating NHP
 CC expression and to produce genetically engineered host cells to express
 CC NHP products in vivo. NHP nucleotide sequences may also be used as part

CC of ribozyme and/or tripe helix sequences that are useful for NHP gene
CC regulation. The NHP polypeptides are useful for generating antibodies, as
CC reagents in diagnostic assays, for identifying other cellular gene
CC products related to NHP and as reagents in assays for screening for
CC compounds that are useful in the treatment of mental, biological or
CC medical disorders and diseases
SQ Sequence 2009 AA;

Query Match 100.0%; Score 28; DB 5; Length 2009;
Best Local Similarity 100.0%; Pred. No. 1.9e-18;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 YMIFVLVIFLGSFYLINILAVVAMAY 28
DB 399 YMIFVLVIFLGSFYLINILAVVAMAY 426

RESULT 25

ABG69292 standard; protein; 2009 AA.

AC ABG69292;

DT 21-OCT-2002 (first entry)

DE Human sodium channel alpha 1-subunit (SCNA1) variant protein #4.

KW Human; sodium channel alpha 1-subunit; SCN1A; episodic ataxia; epilepsy;
KW generalised epilepsy with febrile seizures plus; myasthenia;
KW sodium channel dysfunction; malignant hyperthermia; neuropathic pain;
KW inflammatory pain; Alzheimer's disease; Parkinson's disease; myotonia;
KW schizophrenia; hyperkplexia; anticonvulsant; analgesic; neuroprotective;
KW nootropic; anti-Parkinsonian; neuroleptic.

OS Homo sapiens.

PN W0200250096-A1.

PD 27-JUN-2002.

PF 20-DEC-2001; 2001WO-AU001648.

PR 20-DEC-2000; 2000AU-00002203.

PA (BION-) BIONOMICS LTD.

PI Wallace RH, Mulley JC, Berkovic SF;

DR WPI; 2002-528445/56.

PT New nucleic acid encoding mutant alpha subunit of a mammalian voltage-
PT gated sodium channel, useful for diagnosis of epilepsy, particularly
PT generalized epilepsy with febrile seizures plus.
XX

PS Claim 53; Page 147-157; 198pp; English.

XX The invention relates to a nucleic acid molecule encoding a mutant alpha
CC subunit of a mammalian voltage-gated sodium channel. The DNA and the
CC polypeptide may be used in the diagnosis of epilepsy, in particular
CC generalised epilepsy with febrile seizures plus, and other disorders
CC associated with sodium channel dysfunction. The polypeptide is useful for
CC screening of candidate pharmaceutical agents, where high throughput
CC manufacturing techniques are employed. The sequences are also useful in the
CC manufacture of a medicament for the treatment of a disorder associated
CC with sodium channel dysfunction such as epilepsy, particularly
CC generalised epilepsy with febrile seizures plus, malignant hyperthermia,
CC myasthenia, episodic ataxia, neuropathic and inflammatory pain,
CC Alzheimer's disease, Parkinson's disease, schizophrenia, hyperkplexia
CC and myotonia. Sequences ABG69289-ABG69293 represent human sodium channel
CC alpha 1-subunit (SCNA1) polypeptides of the invention
SQ Sequence 2009 AA;

Query Match 100.0%; Score 28; DB 5; Length 2009;
Best Local Similarity 100.0%; Pred. No. 1.9e-18;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 YMIFVLVIFLGSFYLINILAVVAMAY 28
DB 399 YMIFVLVIFLGSFYLINILAVVAMAY 426

Search completed: January 27, 2005, 17:45:13
Job time : 90.5 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM protein - protein search, using sw model

Run on: January 27, 2005, 17:36:05 ; Search time 17 Seconds
(without alignments)
158.475 Million cell updates/sec

Title: US-10-608-584-1
Perfect score: 28
Sequence: 1 YMIFVTVIFLGSFYLINILAVAMAY 28

Scoring table: OLIGO
Gapop 60.0 , Gapext 60.0

Searched: 283416 seqs, 96216763 residues

Word size : 0

Total number of hits satisfying chosen parameters: 283416

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Listing first 100 summaries

Database :
1: pir1:*
2: pir2:*
3: pir3:*
4: pir4:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	28	100.0	2005	2 A46269	sodium channel alp
2	28	100.0	2005	2 B25019	sodium channel pro
3	28	100.0	2009	2 A25019	sodium channel pro
4	21	75.0	1820	1 CHEB	sodium channel alp
5	21	75.0	1977	2 S54771	sodium channel alp
6	21	75.0	2049	2 T43161	sodium channel pro
7	20	71.4	1835	2 T54323	sodium channel alp
8	20	71.4	1836	2 T54893	sodium channel alp
9	20	71.4	1836	2 J50648	sodium channel alp
10	20	71.4	1836	2 I51964	sodium channel alp
11	20	71.4	1840	1 CHRTM1	sodium channel pro
12	16	57.1	1951	2 S00320	sodium channel pro
13	16	57.1	1983	2 A60054	sodium channel pro
14	13	46.4	1957	2 S68453	sodium channel pro
15	13	39.3	1976	2 I56555	sodium channel pro
16	11	39.3	2016	2 A38195	sodium channel pro
17	11	39.3	2019	2 A33996	sodium channel pro
18	10	35.7	1784	2 T43167	sodium channel pro
19	9	32.1	286	2 S16294	chlorophyll a/b-bi
20	9	32.1	286	2 S21386	chlorophyll a/b-bi
21	9	32.1	1765	2 T42388	sodium channel alp
22	8	28.6	103	2 A53461	voltage-sensitive
23	8	28.6	245	2 T15794	hypothetical prote
24	8	28.6	575	2 D69611	ABC transporter re
25	8	28.6	1034	2 S60051	sodium channel alp
26	8	28.6	1034	2 S60060	sodium channel alp
27	8	28.6	1321	2 A60165	sodium channel pro
28	8	28.6	1689	2 S72467	sodium channel pro
29	8	28.6	1820	2 A33299	sodium channel pro

30	8	28.6	2108	2 S72458	sodium channel pro
31	7	25.0	211	2 F70029	conserved hypothet
32	7	25.0	308	2 J23253	chitinase (EC 3.2.
33	7	25.0	319	2 S14948	chitinase (EC 3.2.
34	7	25.0	349	2 A72605	probable high-affi
35	7	25.0	361	2 T33402	hypothetical prote
36	7	25.0	365	1 S77076	phospho-N-acetylmu
37	7	25.0	376	2 A36978	MAP kinase mpk-1 (
38	7	25.0	444	2 A36977	MAP kinase sur-1 (
39	7	25.0	724	2 B85045	probable calcium c
40	7	25.0	800	2 T26583	hypothetical prote
41	7	25.0	844	2 B69000	cation-transportin
42	7	25.0	1217	2 C86159	hypothetical prote
43	6	21.4	50	2 D91083	hypothetical prote
44	6	21.4	51	2 I61238	heparin-binding ep
45	6	21.4	59	2 S56139	membrane protein n
46	6	21.4	72	2 AH0039	probable exported
47	6	21.4	82	2 B43702	X'82 protein - Afr
48	6	21.4	84	2 S27162	heparin-binding ep
49	6	21.4	103	1 BVMCS	chaperonin groES -
50	6	21.4	103	2 A36721	groES protein - Sy
51	6	21.4	106	2 D69931	hypothetical prote
52	6	21.4	106	2 T22069	hypothetical prote
53	6	21.4	115	2 T17085	NADH2 dehydrogenas
54	6	21.4	118	2 I45348	V18 protein - Afr
55	6	21.4	121	2 A36821	LIS121-1 protein -
56	6	21.4	124	2 B36821	LIS124-2 protein -
57	6	21.4	124	2 C36821	LIS124-1 protein -
58	6	21.4	124	2 C43702	U124 protein - Afr
59	6	21.4	135	2 T05909	membrane protein Y
60	6	21.4	141	2 A86417	probable auxin-ind
61	6	21.4	157	2 D83742	cytochrome c oxida
62	6	21.4	159	2 B70635	hypothetical prote
63	6	21.4	166	2 D70553	hypothetical prote
64	6	21.4	181	2 B84121	hypothetical prote
65	6	21.4	189	2 T02423	probable low tempe
66	6	21.4	197	2 T07995	ycf4 protein - Chl
67	6	21.4	201	2 G72663	hypothetical prote
68	6	21.4	201	2 G90385	hypothetical prote
69	6	21.4	208	1 A38432	heparin-binding EG
70	6	21.4	208	1 A41914	diphtheria toxin re
71	6	21.4	213	2 A75218	hypothetical prote
72	6	21.4	215	2 I38473	olfactory receptor
73	6	21.4	218	2 G95016	conserved hypothet
74	6	21.4	227	2 B81054	deda protein, prob
75	6	21.4	227	2 B70790	hypothetical prote
76	6	21.4	234	2 A70131	phosphatidytranst
77	6	21.4	238	2 T33550	hypothetical prote
78	6	21.4	240	2 H97783	hypothetical prote
79	6	21.4	246	2 S15378	H+-transporting tw
80	6	21.4	247	2 T11315	ATP synthase chain
81	6	21.4	249	2 T27019	hypothetical prote
82	6	21.4	252	2 B98204	probable permease
83	6	21.4	252	2 AF3082	hypothetical prote
84	6	21.4	254	1 G69878	conserved hypothet
85	6	21.4	259	2 B83419	conserved hypothet
86	6	21.4	261	2 B81823	Deda-family' integr
87	6	21.4	261	2 B90107	putative CCR4-asso
88	6	21.4	266	2 C87367	flagellar biosynth
89	6	21.4	273	2 T37841	probable transloca
90	6	21.4	273	2 A11864	hypothetical prote
91	6	21.4	276	2 A72276	phosphate ABC tran
92	6	21.4	279	2 D89784	hypothetical prote
93	6	21.4	280	2 D84015	maltose/maltodextr
94	6	21.4	282	2 A10948	hypothetical prote
95	6	21.4	283	2 E83902	maltose/maltodextr
96	6	21.4	284	2 AC2551	hypothetical prote
97	6	21.4	284	2 E97054	spontaneous protei
98	6	21.4	288	2 T37395	probable 33.6k pro
99	6	21.4	288	2 A42518	A8r protein - vacc
100	6	21.4	288	2 F72164	A8r protein - vari

6 i
1

Query Match	100.0%;	Score 28;	DB 2;	Length 2005;
Best Local Similarity	100.0%;	Pred. No. 2.5e-19;		
Matches 28;	Conservative 0;	Mismatches 0;		

[illegible]

RESULT 2

A/Residues: 1-2005 <NOD>
 A/Cross-references: UNIPROT:Q63509
 A/Experimental source: brain

dition: Development

Query Match	100.0%;	Score
Best Local Similarity	100.0%;	Pr
Matches 28: Consensus		

[illegible]

SULT 3

value protein

1

C/Species: Rattus norvegicus (Norway rat)
C/Date: 30-Jun-1988 #sequence revision 30-Jun-1998 #text_change 09-Jul-2004
C/Accession: A25019; S40783; I84764
R:/Node: M.; Ikeda, T.; Kayano, T.; Suzuki, H.; Takeshima, H.; Kurasaki, M.; Takahashi, H.; Nature 320, 188-197, 1986
A:/Title: Existence of distinct sodium channel messenger RNAs in rat brain.
A:/Reference number: A93377; MUID:86146901; PMID:3754035
A/Accession: A25019
A/Molecule type: mRNA
A/Residues: 1-2009 <NOD>
A/Cross-references: UNIPROT:P04774; GB:X03638; NID:g57216; PID:CAA27886.1; PID:g57217
R:/Experimental source: Brain
R:/Sarcos, R.; Gupta, S.K.; Auld, V.J.; Dunn, R.J.
Nucleic Acids Res. 19, 5673-5679, 1991
A:/Title: Developmentally regulated alternative RNA splicing of rat brain sodium channel
A:/Reference number: S40782; MUID:92051314; PMID:1658739
A/Accession: S40783
A/Molecule type: DNA
A/Residues: 177-253 <SAR>
R:/Node: M.; Numa, S.
J. Recept. Res. 7, 467-497, 1987
A:/Title: Structure and function of sodium channel.
A:/Reference number: I50536; MUID:87311395; PMID:2442385
A/Accession: I84764
A/Status: preliminary; translated from GB/EMBL/DBJ
A/Molecule type: mRNA
A/Residues: 1-2009 <RES>
A/Cross-references: GB:M22253; NID:g1041088; PID:AAA79965.1; PID:g1041089
C/Keyword: duplication; ion transport; sodium channel; transmembrane protein; voltage-

Query Match 100.0%; Score 28; DB 2; Length 2009;
Best Local Similarity 100.0%; Pred. No. 2.5e-19;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 YMFVVLVIFGSPYLNLILAVVAVAY 28
Db 399 YMFVVLVIFGSPYLNLILAVVAVAY 426

RESULT 4
CHER
sodium channel protein - electric eel
C/Species: Electrophorus electricus (electric eel)
C/Date: 28-May-1986 #sequence revision 28-May-1986 #text_change 09-Jul-2004
C/Accession: A03178; I50536
R:/Node: M.; Shimizu, S.; Tanabe, T.; Takai, T.; Kayano, T.; Ikeda, T.; Takahashi, H.; Nature 312, 121-127, 1984
A:/Title: Primary structure of Electrophorus electricus sodium channel deduced from cDNA
A/Reference number: A03178; MUID:85061498; PMID:6205577
A/Accession: A03178
A/Molecule type: mRNA
A/Residues: 1-1820 <NOD>
A/Cross-references: UNIPROT:P02719; GB:X01119; NID:g62776; PID:CAA25587.1; PID:g62777
R:/Node: M.; Numa, S.
J. Recept. Res. 7, 467-497, 1987
A:/Title: Structure and function of sodium channel.
A:/Reference number: I50536; MUID:87311395; PMID:2442385
A/Accession: I50536
A/Status: preliminary; translated from GB/EMBL/DBJ
A/Molecule type: mRNA
A/Residues: 1-1820 <NO2>
A/Cross-references: GB:M22252; NID:g1041048; PID:AAA79960.1; PID:g1041049
C/Comment: This membrane glycoprotein mediates the voltage-dependent sodium-ion permeability of the membrane, the protein forms a sodium-selective channel through which sodium ions pass. This sequence contains four highly homologous internal repeats (excluding repeats 1 and 2) and has a net positive charge (S4), and one is neutral (S2).
C/Comment: The four repeating units are thought to be oriented pseudosymmetrically across the membrane.
C/Comment: The presence of four homologous structures within this molecule is consistent with the possibility that activation and inactivation gates are located near the extracellular surface.
C/Comment: Available data suggest that positively charged residues of S4, act as a voltage sensor.

C:Superfamily: sodium channel protein
C:Keywords: duplication; glycoprotein; ion transport; membrane protein; sodium channel;
F:111-141,555-807,989-1281,1311-1341/Region: duplication internal repeats I, II, III and
F:111-141,555-585,989-1019,1311-1341/Region: S1 of repeats I through IV
F:150-171,597-620,1033-1057,1153-1376/Region: S2 of repeats I through IV
F:177-197,628-643,1062-1079,1381-1398/Region: S3 of repeats I through IV
F:204-224,651-671,1092-1112,1417-1437/Region: S4 of repeats I through IV
F:244-264,691-711,1132-1152,1454-1474/Region: S5 of repeats I through IV
F:379-402,767-790,1236-1264,1544-1567/Region: S6 of repeats I through IV
F:205,278,288,317,591,690,797,1160,1174,1806/Binding site: carbohydrate (Asn) (covalent)

Query Match 75.0%; Score 21; DB 1; Length 1820;
Best Local Similarity 100.0%; Pred. No. 1.5e-12;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 VIFLGSFYILNLIILAVVAMAY 28
|||||
DB 385 VIFLGSFYILNLIILAVVAMAY 405

RESULT 5
S54771
sodium channel alpha subunit - human
C:Species: Homo sapiens (man)
C:Date: 27-Oct-1995 #sequence_revision 03-Nov-1995 #text_change 09-Jul-2004
C:Accession: S54771
R:Kluehauer, N.; Iacino, L.; Flockerzi, V.; Hofmann, F.
EMBO J. 14, 1084-1090, 1995
A:Title: Structure and functional expression of a new member of the tetrodotoxin-sensitive
A:Reference number: S54771; PMID:95237189; PMID:7720699
A:Accession: S54771
A:Status: preliminary; nucleic acid sequence not shown
A:Molecule type: mRNA
A:Residues: 1-197 <KLU>
A:Cross-references: UNIPROT:Q15858, EMBL:X62835, NID:G758109, PIDN:CAA58042.1; PID:G7581
C:Superfamily: sodium channel protein
C:Keywords: duplication

Query Match 75.0%; Score 21; DB 2; Length 1977;
Best Local Similarity 100.0%; Pred. No. 1.6e-12;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 VIFLGSFYILNLIILAVVAMAY 28
|||||
DB 385 VIFLGSFYILNLIILAVVAMAY 405

RESULT 6
T43161
sodium channel protein Tuna1 - sea squirt (Halocynthia roretzi)
C:Species: Halocynthia roretzi
C:Date: 11-Jan-2000 #sequence_revision 11-Jan-2000 #text_change 09-Jul-2004
C:Accession: T43161
R:Okamura, Y.; Ono, F.; Okagaki, R.; Chong, J.; Mandel, G.
Neuron 13, 937-948, 1994
A:Title: Neutral expression of a sodium channel gene requires cell-specific interactions.
A:Reference number: Z2320; PMID:95033215; PMID:794638
A:Accession: T43161
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-2049 <OKA>
A:Cross-references: UNIPROT:Q25150; EMBL:DJ7311; PIDN:BA04133.1
C:Superfamily: sodium channel protein
C:Keywords: sodium channel; transmembrane protein

Query Match 75.0%; Score 21; DB 2; Length 2049;
Best Local Similarity 100.0%; Pred. No. 1.6e-12;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 VIFLGSFYILNLIILAVVAMAY 28
|||||
DB 449 VIFLGSFYILNLIILAVVAMAY 469

RESULT 7
I54323
sodium channel alpha subunit - human
C:Species: Homo sapiens (man)
C:Date: 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change 20-Aug-1999
C:Accession: I54323
R:McClatchey, A.L.; Lin, C.S.; Wang, J.; Hoffman, E.P.; Rojas, C.; Gueelle, J.F.
Hum. Mol. Genet. 1, 521-527, 1992
A:Title: The genomic structure of the human skeletal muscle sodium channel gene.
A:Reference number: I54323; PMID:9335844; PMID:1339144
A:Accession: I54323
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-1835 <RES>
A:Cross-references: GB:I01983; NID:G337992; PIDN:AAA75557.1; PID:G908809
C:Genetics:
C:Gene: GDB:SCN4A
A:Cross-references: GDB:125181; OMIM:170500
A:Map position: 17q23.1-17q25.3
A:Introns: 91/3; 131/2; 161/2; 204/2; 235/1; 346/1; 367/2; 414/3; 484/3; 536/1; 615/3;
C:Superfamily: sodium channel protein
C:Keywords: duplication

Query Match 71.4%; Score 20; DB 2; Length 1835;
Best Local Similarity 100.0%; Pred. No. 1.4e-11;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 9 IFLGSFYILNLIILAVVAMAY 28
|||||
DB 431 IFLGSFYILNLIILAVVAMAY 450

RESULT 8
I64893
sodium channel alpha subunit - human
C:Species: Homo sapiens (man)
C:Date: 06-Sep-1996 #sequence_revision 06-Sep-1996 #text_change 09-Jul-2004
C:Accession: I64893
R:George, A.L.
Ann. Neurol. 31, 131-137, 1992
A:Title: Primary structure of the adult human skeletal muscle voltage-dependent sodium
A:Reference number: I51964; PMID:92246457; PMID:115496
A:Accession: I64893
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-1836 <RES>
A:Cross-references: UNIPROT:P35499; GB:W81758; NID:G338212; PIDN:AAA6054.1; PID:G33821
A:Genetics:
A:Gene: SKM1
C:Superfamily: sodium channel protein
C:Keywords: duplication

Query Match 71.4%; Score 20; DB 2; Length 1836;
Best Local Similarity 100.0%; Pred. No. 1.4e-11;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 9 IFLGSFYILNLIILAVVAMAY 28
|||||
DB 431 IFLGSFYILNLIILAVVAMAY 450

RESULT 9
J50648
sodium channel alpha chain - human
C:Species: Homo sapiens (man)
C:Date: 30-Jun-1992 #sequence_revision 30-Jun-1992 #text_change 09-Jul-2004
C:Accession: J50648; J42099
R:Yang, J.; Rojas, C.V.; Zhou, J.; Schwartz, L.S.; Nicholas, H.; Hofmann, E.P.
Biochem. Biophys. Res. Commun. 182, 794-801, 1992
A:Title: Sequence and genomic structure of the human adult skeletal muscle sodium channe
A:Reference number: J50648; PMID:92134303; PMID:1310396
A:Accession: J50648

A>Status: nucleic acid sequence not shown
A:Molecule type: mRNA
A:Residues: 1-1836 <MAN>
A:Cross-references: UNIPROT:P35499
A>Note: 861-Ap was also found as the result of polymorphism
R:McClatchey, A.L.; Van den Berg, P.; Perleak-Vance, M.A.; Raskind, W.; Verelien, C.; M
Cell 66, 769-774, 1992
A>Title: Temperature-sensitive mutations in the III-IV cytoplasmic loop region of the sk
A:Reference number: A42099; MUID:922454689; PMID:1310898
A:Accession: A42099
A:Molecule type: DNA
A:Residues: 1299-1351 <MCC>
A:Cross-references: GB:S82622; NID:9245611; PIDN:AAB21450.1; PID:9245612
A:Experimental source: skeletal muscle
A>Note: sequence extracted from NCBI backbone (NCBI:82622, NCBI:82623)
C:Genetics:
A:Gene: GDB:SCN4A
A:Cross-references: GDB:125181; OMIM:170500
A:Map position: 17q23.1-17q25.3
C:Superfamily: sodium channel protein
C:Keywords: duplication; glycoprotein; phosphoprotein; transmembrane protein
F:129-150/Domain: transmembrane #status predicted <TR1>
F:159-178/Domain: transmembrane #status predicted <TR2>
F:191-210/Domain: transmembrane #status predicted <TR3>
F:217-236/Domain: transmembrane #status predicted <TR4>
F:253-269/Domain: transmembrane #status predicted <TR5>
F:424-449/Domain: transmembrane #status predicted <TR6>
F:574-597/Domain: transmembrane #status predicted <TR7>
F:609-632/Domain: transmembrane #status predicted <R11>
F:641-660/Domain: transmembrane #status predicted <R12>
F:667-686/Domain: transmembrane #status predicted <R13>
F:702-724/Domain: transmembrane #status predicted <R14>
F:777-802/Domain: transmembrane #status predicted <R15>
F:1027-1049/Domain: transmembrane #status predicted <R16>
F:1064-1089/Domain: transmembrane #status predicted <R17>
F:1096-1116/Domain: transmembrane #status predicted <R18>
F:1122-1143/Domain: transmembrane #status predicted <R19>
F:1163-1184/Domain: transmembrane #status predicted <R20>
F:1262-1285/Domain: transmembrane #status predicted <R21>
F:1349-1372/Domain: transmembrane #status predicted <R22>
F:1384-1407/Domain: transmembrane #status predicted <R23>
F:1441-1437/Domain: transmembrane #status predicted <R24>
F:1447-1469/Domain: transmembrane #status predicted <R25>
F:1485-1507/Domain: transmembrane #status predicted <R26>
F:1574-1598/Domain: transmembrane #status predicted <R27>
F:21-149, 220, 378, 415, 1019, 1130, 1242, 1313, 1721, 1826/Binding site: phosphate (Thr) (coval
F:56, 251, 513, 653, 1511, 1746/Binding site: phosphate (Ser) (covalent) (by protein kinase A
F:214, 288, 291, 297, 303, 315, 321, 333, 362, 507, 702, 961, 1191, 1205/Binding site: phosphate (Ser) (covalent) (by protei
F:214, 288, 291, 297, 303, 315, 321, 333, 362, 507, 702, 961, 1191, 1205/Binding site: phosphate (Ser) (covalent) (by protei
F:387, 457/Binding site: phosphate (Thr) (covalent) (by protein kinase A) #status predict

Query Match
Best Local Similarity 71.4%; Score 20; DB 2; Length 1836;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 9 IFGSSFYLLNLIILAVVAVMAY 28
431 IFGSSFYLLNLIILAVVAVMAY 450

RESULT 10
151964
sodium channel alpha chain SCN4A, skeletal muscle - human
C:Species: Homo sapiens (man)
C>Date: 24-May-1996 #sequence_revision 24-May-1996 #text_change 09-Jul-2004
R:Georgie, A.L.
Ann. Neurol. 31, 131-137, 1992
A>Title: Primary structure of the adult human skeletal muscle voltage-dependent sodium
A:Reference number: 151964; MUID:92246457; PMID:1315496
A:Accession: 151964
A>Status: preliminary; translated from GB/EMBL/DBD

A:Residues: 1-1836 <RES>
A:Cross-references: UNIPROT:P35499; GB:L04236; NID:9292485; PIDN:AAB59624.1; PID:9292487
C:Genetics:
A:Gene: GDB:SCN4A
A:Cross-references: GDB:125181; OMIM:170500
A:Map position: 17q23.1-17q25.3
A:Introns: 91/3; 131/2; 161/2; 204/2; 235/1; 346/1; 367/2; 414/3; 484/3; 536/1; 615/3;
C:Superfamily: sodium channel protein
C:Keywords: duplication; skeletal muscle

Query Match
Best Local Similarity 71.4%; Score 20; DB 2; Length 1836;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 9 IFGSSFYLLNLIILAVVAVMAY 28
431 IFGSSFYLLNLIILAVVAVMAY 450

RESULT 11

CHR1M1
sodium channel protein mul alpha chain, skeletal muscle - rat
C:Species: Rattus norvegicus (Norway rat)
C>Date: 30-Sep-1990 #sequence_revision 30-Sep-1990 #text_change 09-Jul-2004
C:Accession: JN0007
R:Titmer, J.S.; Cooperman, S.S.; Tomiko, S.A.; Zhou, J.; Crean, S.M.; Boyle, M.B.; Kal
Neuron 3, 33-49, 1989
A>Title: Primary structure and functional expression of a mammalian skeletal muscle sod
A:Reference number: JN0007; MUID:90148778; PMID:2559760
A:Accession: JN0007
A:Molecule type: mRNA
A:Residues: 1-1840 <R1>
A:Cross-references: UNIPROT:PI5390; GB:M26643; NID:9205651; PIDN:AAA1662.1; PID:9205652
C:Comment: Action potentials propagated by skeletal muscle sodium channels are responsib
C:Comment: This heavily glycosylated protein contains four homologous domains, each of w
C:Superfamily: sodium channel protein
C:Keywords: duplication; glycoprotein; ion transport; neuromuscular junction; phosphopro
F:120-458, 561-813, 1013-1305, 1335-1611/Region: duplication
F:56, 251, 1221, 1504/Binding site: phosphate (Ser) (covalent) (by CAMP-dependent kinase) #
F:214, 288, 291, 297, 303, 309, 315, 327, 356, 502, 696, 954, 1184, 1198, 1563, 1702/Binding site: carb

Query Match
Best Local Similarity 71.4%; Score 20; DB 1; Length 1840;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 9 IFGSSFYLLNLIILAVVAVMAY 28
425 IFGSSFYLLNLIILAVVAVMAY 444

RESULT 12

S00320
sodium channel protein III - rat
C:Species: Rattus norvegicus (Norway rat)
C>Date: 30-Jun-1989 #sequence_revision 30-Jun-1989 #text_change 09-Jul-2004
C:Accession: S00320
R:Kayano, T.; Noda, M.; Flockerzi, V.; Takahashi, H.; Numa, S.
FEBS Lett. 228, 187-194, 1988
A>Title: Primary structure of rat brain sodium channel III deduced from the cDNA sequenc
A:Reference number: S00320; MUID:8813594; PMID:2449363
A:Accession: S00320
A:Molecule type: mRNA
A:Residues: 1-1951 <RAY>
A:Cross-references: UNIPROT:P08104; EMBL:Y00766; NID:957210; PIDN:CAA68735.1; PID:957211
A>Note: 270-1le, 278-1eu, 355-Thr, 513-Lys, and 1059-Arg were also found
C:Superfamily: sodium channel protein
C:Keywords: duplication; transmembrane protein

Query Match
Best Local Similarity 57.1%; Score 16; DB 2; Length 1951;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 YMIFFVLVIFLGSFYL 16
 Db 400 YMIFFVLVIFLGSFYL 415

RESULT 13

A60054

sodium channel protein IIB, long form - rat
 C:Species: Rattus norvegicus (Norway rat)

C>Date: 03-Mar-1993 #sequence_revision 03-Mar-1993 #text_change 09-Jul-2004

C:Accession: A60054; B44824

C:Joho, R.H.; Moorman, J.R.; Vandongen, A.M.J.; Kirsch, G.E.; Silberberg, H.; Schuster, Brain Res. Mol. Brain Res. 7, 105-113, 1990

A>Title: Toxin and kinetic profile of rat brain type III sodium channels expressed in X

A:Reference number: A60054; MUID:90251117; PMID:2160038

A:Accession: A60054

A:Status: not compared with conceptual translation

A:Molecule type: mRNA

A:Residues: 1-1983 <JOH>

A:Cross-references: UNIPROT:064243

R:Schaller, K.L.; Krzemien, D.M.; McKenna, N.M.; Caldwell, J.H.

J. Neurosci. 12, 1370-1381, 1992

A>Title: Alternatively spliced sodium channel transcripts in brain and muscle.

A:Reference number: A44824; MUID:92211397; PMID:1313493

A:Accession: B44824

A:Status: preliminary

A:Molecule type: mRNA

A:Residues: 611-662 <SCH>

A:Cross-references: GB:S97388; NID:G248225; PIDN:AA821984.1; PID:G248226

A:Experimental source: skeletal muscle

A>Note: sequence inconsistent with the nucleotide translation

C:Superfamily: sodium channel protein

C:Keywords: duplication; glycoprotein; ion transport; sodium channel; transmembrane proc

Query Match 57.1%; Score 16; DB 2; Length 1983;

Best Local Similarity 100.0%; Pred. No. 1.2e-07;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 YMIFFVLVIFLGSFYL 16

Db 400 YMIFFVLVIFLGSFYL 415

RESULT 14

sodium channel protein SNS - rat

C:Species: Rattus norvegicus (Norway rat)

C>Date: 17-Jul-1998 #sequence_revision 17-Jul-1998 #text_change 09-Jul-2004

C:Accession: S68453

R:Akopian, A.N.; Sivijott, L.; Wood, J.N.

Nature 379, 257-262, 1996

A>Title: A tetrodotoxin-resistant voltage-gated sodium channel expressed by sensory neu

A:Reference number: S68453; MUID:96138382; PMID:8538791

A:Accession: S68453

A:Status: nucleic acid sequence not shown

A:Molecule type: mRNA

A:Residues: 1-1957 <AKO>

A:Cross-references: UNIPROT:063554; GB:X92184; NID:G1209466; PIDN:CAA63095.1; PID:G12094

A:Experimental source: dorsal root ganglia

C:Superfamily: sodium channel protein

C:Keywords: sodium channel; transmembrane protein; voltage-gated ion channel

F:113-148/Domain: transmembrane #status predicted <TM1>

F:158-174/Domain: transmembrane #status predicted <TM2>

F:123-241/Domain: transmembrane #status predicted <TM3>

F:124-265/Domain: transmembrane #status predicted <TM4>

F:376-392/Domain: transmembrane #status predicted <TM5>

F:666-682/Domain: transmembrane #status predicted <TM6>

F:702-718/Domain: transmembrane #status predicted <TM7>

F:731-747/Domain: transmembrane #status predicted <TM8>

F:788-804/Domain: transmembrane #status predicted <TM9>

F:865-881/Domain: transmembrane #status predicted <TM10>

F:1156-1172/Domain: transmembrane #status predicted <TM11>

F:1194-1210/Domain: transmembrane #status predicted <TM12>

F:1221-1237/Domain: transmembrane #status predicted <TM13>

F:1286-1302/Domain: transmembrane #status predicted <TM14>

F:1400-1416/Domain: transmembrane #status predicted <TM15>

F:1482-1498/Domain: transmembrane #status predicted <TM16>

F:1516-1532/Domain: transmembrane #status predicted <TM17>

F:1546-1562/Domain: transmembrane #status predicted <TM18>

F:1606-1622/Domain: transmembrane #status predicted <TM19>

F:1708-1724/Domain: transmembrane #status predicted <TM20>

Query Match 46.4%; Score 13; DB 2; Length 1957;

Best Local Similarity 100.0%; Pred. No. 9.6e-05;

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 FPLVIFLGSFYL 16

Db 375 FPLVIFLGSFYL 387

RESULT 15

sodium channel protein 6 - rat

C:Species: Rattus norvegicus (Norway rat)

C>Date: 26-Jul-1996 #sequence_revision 26-Jul-1996 #text_change 09-Jul-2004

C:Accession: I56555

R:Schaller, K.L.; Krzemien, D.M.; Yarowsky, P.J.; Krueger, B.K.; Caldwell, J.H.

J. Neurosci. 15, 3231-3242, 1995

A>Title: A novel, abundant sodium channel expressed in neurons and glia.

A:Reference number: I56555; MUID:95271284; PMID:7751906

A:Accession: I56555

A:Status: preliminary; translated from GB/EMBL/DBJ

A:Molecule type: mRNA

A:Residues: 1-1976 <RES>

A:Cross-references: UNIPROT:063541; GB:L39018; NID:9829033; PIDN:AA642059.1; PID:982903

C:Genetic:

A:Gene: SCP6

C:Superfamily: sodium channel protein

C:Keywords: duplication

Query Match 39.3%; Score 11; DB 2; Length 1976;

Best Local Similarity 100.0%; Pred. No. 0.0086;

Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 18 NLIILAVVAMAY 28

Db 405 NLIILAVVAMAY 415

RESULT 16

sodium channel protein hH1, cardiac - human

N:Alternate names: tetrodotoxin-insensitive, voltage-dependent sodium channel, TTX-I Na

C:Species: Homo sapiens (man)

C>Date: 31-Dec-1993 #sequence_revision 31-Dec-1993 #text_change 09-Jul-2004

C:Accession: A38195

R:Cellen, M.E.; George Jr., A.L.; Chen, L.Q.; Chahine, M.; Horn, R.; Barchi, R.L.; Kai

Proc. Natl. Acad. Sci. U.S.A. 89, 554-558, 1992

A>Title: Primary structure and functional expression of the human cardiac tetrodotoxin-

A:Reference number: A38195; MUID:92115699; PMID:1309946

A:Accession: A38195

A:Status: nucleic acid sequence not shown

A:Molecule type: mRNA

A:Residues: 1-2016 <GEU>

A:Cross-references: UNIPROT:Q14524; GB:M77235; NID:G184038; PIDN:AAA58644.1; PID:G18403

A:Experimental source: heart

C:Superfamily: sodium channel protein

C:Keywords: cardiac muscle; duplication; glycoprotein; heart; ion transport; sodium char

Query Match 39.3%; Score 11; DB 2; Length 2016;

Best Local Similarity 100.0%; Pred. No. 0.0087;

Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 18 NLIILAVVAMAY 28

Db 406 NLILAVVAMAY 416

RESULT 17

sodium channel protein I, cardiac - rat
A:Accession: A33996
N:Alternate names: sodium channel protein (SKM2) alpha chain
C:Species: Rattus norvegicus (Norway rat)
C:Date: 30-Mar-1990 #sequence_revision 30-Mar-1990 #text_change 09-Jul-2004

C:Accession: A33996; U00412
R:Rogart, R.B.; Cribbs, L.L.; Muglia, L.K.; Kephart, D.D.; Kaiser, M.W.
Proc. Natl. Acad. Sci. U.S.A. 86, 8170-8174, 1989

A:Title: Molecular cloning of a putative tetrodotoxin-resistant rat heart Na(+) channel
A:Reference number: A33996; PMID:90046760; PMID:25543302
A:Accession: A33996

A:Status: preliminary
A:Molecule type: mRNA

A:Residues: 1-2019 <ROG>
A:Cross-references: UNIPROT:P15389; GB:M27902; NID:9206857; PIDN:AAA42114.1; PID:9206858
R:Kallen, R.G.; Sheng, Z.H.; Yang, J.; Chen, L.; Rogart, R.B.; Barchi, R.L.
Neuron 4, 233-242, 1990

A:Title: Primary structure and expression of a sodium channel characteristic of denervat
A:Reference number: U00412; PMID:9016613; PMID:2155010
A:Accession: U00412

A:Molecule type: mRNA
A:Residues: 1-479, 481-1712, 'T', 1714-1963, 'R', 1965-2019 <KAL>
A:Experimental source: muscle
C:Superfamily: sodium channel protein

C:Keywords: cardiac muscle; duplication; heart; sodium channel; transmembrane protein

Query Match 39.3%; Score 11; DB 2; Length 2019;
Best Local Similarity 100.0%; Pred. No. 0.0087;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 18 NLILAVVAMAY 28
Db 407 NLILAVVAMAY 417

RESULT 18

sodium channel protein - California market squid
C:Species: Loligo opalescens (California market squid)
C:Date: 11-Jan-2000 #sequence_revision 11-Jan-2000 #text_change 09-Jul-2004

C:Accession: T43167
R:Rosenthal, J.J.; Gilly, W.F.
Proc. Natl. Acad. Sci. U.S.A. 90, 10026-10030, 1993

A:Title: Amino acid sequence of a putative sodium channel expressed in the giant axon of
A:Reference number: Z22324; PMID:94052096; PMID:8234251
A:Accession: T43167

A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-1784 <ROS>

A:Cross-references: UNIPROT:Q25377; EMBL:L1979; NID:9349118; PID:9349119; PIDN:AAA16202
A:Experimental source: stellate ganglia
C:Superfamily: sodium channel protein

C:Keywords: ion transport; membrane protein; sodium channel; voltage-gated ion channel

Query Match 35.7%; Score 10; DB 2; Length 1784;
Best Local Similarity 100.0%; Pred. No. 0.074;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 7 LVIFLGSFYL 16
Db 383 LVIFLGSFYL 392

RESULT 19

chlorophyll a/b-binding protein type I precursor - tomato
C:Species: Lycopersicon esculentum (tomato)
C:Date: 19-Mar-1997 #sequence_revision 19-Mar-1997 #text_change 09-Jul-2004

C:Accession: S16294
R:Picchersky, E.; Subramaniam, R.; White, M.J.; Reid, J.; Aebersold, R.; Green, B.R.
Mol. Gen. Genet. 227, 277-284, 1991

A:Title: Chlorophyll a/b binding (CAB) polypeptides of CP29, the internal chlorophyll a
or a second CP29 polypeptide.
A:Reference number: S16294; PMID:91287707; PMID:2062308

A:Accession: S16294
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-286 <GRE>

A:Cross-references: UNIPROT:Q00321
C:Superfamily: chlorophyll a/b-binding protein
C:Keywords: chloroplast; thylakoid

Query Match 32.1%; Score 9; DB 2; Length 286;
Best Local Similarity 100.0%; Pred. No. 0.16;
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 17 INILAVVA 25
Db 175 INILAVVA 183

RESULT 20

chlorophyll a/b-binding protein CP29 precursor - barley
C:Species: Hordeum vulgare (barley)
C:Date: 20-Feb-1995 #sequence_revision 20-Feb-1995 #text_change 09-Jul-2004

C:Accession: S21386
R:Sorensen, A.B.; Jensen, B.F.; Gausing, K.
submitted to the EMBL Data Library, October 1991

A:Description: Barley (Hordeum vulgare) gene for CP29, a core chlorophyll a/b binding pr
A:Reference number: S21386
A:Accession: S21386

A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-286 <SOR>

A:Cross-references: UNIPROT:Q40039; EMBL:X63052; NID:918957; PIDN:CAA44777.1; PID:918958
C:Genetics:
A:Introns: 70/1; 98/2; 115/2; 155/3; 200/3
C:Superfamily: chlorophyll a/b-binding protein
C:Keywords: chloroplast; thylakoid

Query Match 32.1%; Score 9; DB 2; Length 286;
Best Local Similarity 100.0%; Pred. No. 0.16;
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 17 INILAVVA 25
Db 175 INILAVVA 183

RESULT 21

sodium channel alpha chain - rat
C:Species: Rattus norvegicus (Norway rat)
C:Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 03-Jul-2004

C:Accession: T42388
R:Diib-Hajj, S.D.; Tyrrell, L.; Black, J.A.; Waxman, S.G.
Proc. Natl. Acad. Sci. U.S.A. 95, 8963-8968, 1998

A:Title: NAN, a novel voltage-gated Na channel, is expressed preferentially in periph
A:Reference number: Z22149; PMID:98338024; PMID:9671787
A:Accession: T42388

A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-1765 <DIB>

A:Cross-references: UNIPROT:O88457; EMBL:AF059030; NID:93372614; PID:93372615; PIDN:AAC4
A:Experimental source: strain Sprague-Dawley; dorsal root ganglia
A:Note: preferentially expressed in sensory neurons within dorsal root ganglia and trig
C:Superfamily: sodium channel protein

Query Match 32.1%; Score 9; DB 2; Length 1765;
Best Local Similarity 100.0%; Pred. No. 0.69;

Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 8 VFVLSFYLL 16
 DB 378 VFVLSFYLL 386

RESULT 22

A53461
 voltage-sensitive sodium channel - house fly (fragment)

C:Species: Musca domestica (house fly)

C:Date: 06-Oct-1994 #sequence_revision 18-Nov-1994 #text_change 09-Jul-2004

C:Accession: A53461

R:Knippl, D.C.; Doyle, K.E.; Marsella-Herrick, P.A.; Soderlund, D.M.

Proc. Natl. Acad. Sci. U.S.A. 91, 2483-2487, 1994

A:Title: Tight genetic linkage between the kdr insecticide resistance trait and a voltage

A:Reference number: A53461; PMID:9419576; PMID:8146143

A:Contents: NAIDM, insecticide-susceptible

A:Accession: A53461

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-103 <KNT>

A:Cross-references: UNIPROT:Q25439, UNIPROT:Q25440, UNIPROT:Q94615

A:Note: sequence inconsistent with nucleotide translation

C:Superfamily: sodium channel protein

C:Keywords: duplication

Query Match 28.6%; Score 8; DB 2; Length 103;
 Best Local Similarity 100.0%; Pred. No. 0.63;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 9 IFVLSFYLL 16
 DB 65 IFVLSFYLL 72

RESULT 23

T15794

hypothetical protein C42D8.1 - Caenorhabditis elegans

C:Species: Caenorhabditis elegans

C:Date: 20-Sep-1999 #sequence_revision 20-Sep-1999 #text_change 09-Jul-2004

C:Accession: T15794

R:Hallsworth, K.

submitted to the EMBL Data Library, April 1996

A:Description: The sequence of C. elegans cosmid C42D8.

A:Reference number: Z18405

A:Accession: T15794

A:Status: preliminary; translated from GB/EMBL/DBJ

A:Molecule type: DNA

A:Residues: 1-245 <HAU>

A:Cross-references: UNIPROT:Q18582, EMBL:U56966, NID:G1293844, PIDN:AAA987

A:Experimental source: strain Bristol N2; clone C42D8

C:Genetics:

A:Gene: CESP:C42D8.1

A:Map position: X

A:Introns: 16/2; 62/3; 90/3; 123/3; 159/3; 208/3

Query Match 28.6%; Score 8; DB 2; Length 245;
 Best Local Similarity 100.0%; Pred. No. 1.3;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 FVLVIFLG 12
 DB 125 FVLVIFLG 132

RESULT 24

D69611

ABC transporter required for expression of cytochrome bd (ATP-) cydD - Bacillus subtilis

C:Species: Bacillus subtilis

C:Date: 05-Dec-1997 #sequence_revision 05-Dec-1997 #text_change 16-Aug-2004

C:Accession: D69611

R:Kunst, F.; Ogasawara, N.; Moszer, I.; Albertini, A.M.; Alloni, G.; Azevedo, V.; Bertei
 C.; Bron, S.; Brouillet, S.; Bruch, C.V.; Caldwell, B.; Capuano, V.; Carter, N.M.; Chk
 A.; Ehrlich, S.D.; Emerson, P.T.; Entian, K.D.; Erington, J.; Fabret, C.; Ferrari, E.
 Nature 390, 249-256, 1997

A:Authors: Foulger, D.; Fritz, C.; Fujita, M.; Fujita, Y.; Fuma, S.; Gallazzi, A.; Galle
 lech, J.; Harwood, C.R.; Henaut, A.; Hilbert, H.; Holtschappel, S.; Hosono, S.; Hullo, M.F
 Koester, P.; Koningsstein, G.; Krogh, S.; Kumano, M.; Kurita, K.; Lapidus, A.; Lardinois
 A:Authors: Lauber, J.; Lazarevic, V.; Lee, S.M.; Levine, A.; Liu, H.; Masuda, S.; Maue
 Y. M.; Ogawa, K.; Ogiwara, A.; Oudega, B.; Park, S.H.; Pario, V.; Pohl, T.M.; Portetelli
 Rieger, M.; Rivolta, C.; Rocha, E.; Roche, B.; Rose, F.; Scalfone, F.; Sekiguchi, Y.; Sekowska, A.; Sero
 A:Authors: Schleich, S.; Schroeter, R.; Scrofano, F.; Sekiguchi, Y.; Sekowska, A.; Sero
 akuch, M.; Tamakoshi, A.; Tanaka, T.; Terpetra, P.; Tognoni, A.; Tosato, V.; Uchiyama,
 T.; Winters, P.; Wipat, A.; Yamamoto, H.; Yamane, K.; Yasumoto, K.; Yata, K.; Yoshida, I
 A:Authors: Yoshikawa, H.F.; Zumestein, E.; Yoshikawa, H.; Danchin, A.

A:Title: The complete genome sequence of the Gram-positive bacterium Bacillus subtilis.

A:Reference number: A63580; PMID:9804403; PMID:9384377

A:Accession: D69611

A:Status: preliminary; nucleic acid sequence not shown; translation not shown

A:Molecule type: DNA

A:Residues: 1-575 <KUN>

A:Cross-references: UNIPROT:P94367; GB:299123; GB:AL009126; NID:G2636240; PIDN:CA15899.

A:Experimental source: strain 168

C:Genetics:

A:Gene: cydD

C:Superfamily: ATP-binding cassette homology

C:Keywords: ATP; nucleotide binding; P-loop

F:353-546/Domain: ATP-binding cassette homology <ABC>

F:370-377/Region: nucleotide-binding motif A (P-loop)

Query Match 28.6%; Score 8; DB 2; Length 575;
 Best Local Similarity 100.0%; Pred. No. 2.6;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 FVLVIFLG 12
 DB 18 FVLVIFLG 25

RESULT 25

S60051

sodium channel alpha chain, exon c-containing splice form - fruit fly (Drosophila virilis)

C:Species: Drosophila virilis

C:Date: 24-Aug-1996 #sequence_revision 08-Nov-1996 #text_change 09-Jul-2004

C:Accession: S60051

R:Thackeray, J.R.; Ganetzky, B.

Genetics 141, 203-214, 1995

A:Title: Conserved alternative splicing patterns and splicing signals in the Drosophila

A:Reference number: S60051; PMID:96042905; PMID:8536968

A:Accession: S60051

A:Status: nucleic acid sequence not shown

A:Molecule type: nucleic acid

A:Residues: 1-1034 <THA>

A:Cross-references: UNIPROT:Q24714; EMBL:U26343

C:Genetics:

A:Gene: FlyBase:FlyBase:FBgn0015214

A:Cross-references: FlyBase:FBgn0015214

C:Superfamily: sodium channel protein

C:Keywords: alternative splicing; duplication; transmembrane protein

F:306-329/Region: alternatively spliced segment 1 (exon 4) #status experimental

F:330-350/Region: alternatively spliced exon a #status experimental

F:338-545/Region: alternatively spliced segment b (exon 9) #status experimental

F:874-886/Region: alternatively spliced segment c (exon 12) #status experimental

F:887-896/Region: alternatively spliced segment d (exon 13) #status experimental

F:958-982/Region: alternatively spliced segment h (exon 14) #status experimental

Query Match 28.6%; Score 8; DB 2; Length 1034;
 Best Local Similarity 100.0%; Pred. No. 4.2;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 9 IFVLSFYLL 16
 DB 188 IFVLSFYLL 195

Fri Jan 28 09:32:02 2005

us-10-608-584-1.011.rpr

Page 8

Search completed: January 27, 2005, 17:52:42
Job time : 20 secs

GenCore version 5.1.6
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OM protein - protein search, using BW model

Run on: January 27, 2005, 17:35:20 ; Search time 92.5 Seconds
(without alignments)
174.167 Million cell updates/sec

Title: US-10-608-584-1
Perfect score: 28
Sequence: 1 YWIFVLVFLGSPYLINLILAVMAY 28

Scoring table: OLIGO
Gapop 60.0 , Gapext 60.0

Searched: 1825181 seqs, 575374646 residues

Word size : 0

Total number of hits satisfying chosen parameters: 1825181

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Listing first 100 summaries

Database : Uniprot_02:*
1: uniprot_sprot:*
2: uniprot_trembl:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	28	100.0	506	2 Q86X25	Q86X25 homo sapien
2	28	100.0	510	2 Q86XY2	Q86XY2 homo sapien
3	28	100.0	1981	2 O81UJ6	O81UJ6 homo sapien
4	28	100.0	2005	1 C1N2_HUMAN	C1N2_HUMAN
5	28	100.0	2005	1 C1N2_RAT	C1N2_RAT
6	28	100.0	2009	1 C1N1_HUMAN	C1N1_HUMAN
7	28	100.0	2009	1 C1N1_RAT	C1N1_RAT
8	21	75.0	213	2 O42420	O42420 gallus gall
9	21	75.0	225	2 O42420	O42420 gallus gall
10	21	75.0	247	2 Q91BD6	Q91BD6 takifugu pa
11	21	75.0	1820	1 C1N4_ELEEL	C1N4_ELEEL
12	21	75.0	1880	2 Q91BF1	Q91BF1 takifugu pa
13	21	75.0	1977	2 Q15858	Q15858 homo sapien
14	21	75.0	1984	2 Q28644	Q28644 cryotolagus
15	21	75.0	1984	2 Q08562	Q08562 rattus norv
16	21	75.0	2049	2 Q25150	Q25150 halocynthia
17	20	71.4	1836	1 C1N4_HUMAN	C1N4_HUMAN
18	20	71.4	1840	1 C1N4_RAT	C1N4_RAT
19	20	71.4	1840	2 Q70611	Q70611 rattus norv
20	20	71.4	1841	2 Q9ER60	Q9ER60 mus musculu
21	16	57.1	530	2 Q90518	Q90518 fuqua rubrip
22	16	57.1	711	2 Q803T8	Q803T8 brachydanio
23	16	57.1	1834	2 Q28371	Q28371 equus caball
24	16	57.1	1951	1 C1N3_RAT	C1N3_RAT
25	16	57.1	1951	2 Q9C007	Q9C007 homo sapien
26	16	57.1	1956	2 Q9Y5Y9	Q9Y5Y9 homo sapien
27	16	57.1	2000	1 C1N3_HUMAN	C1N3_HUMAN
28	16	57.1	2007	2 Q9YGN7	Q9YGN7 cynops pyrr
29	13	46.4	1956	2 Q62968	Q62968 rattus norv
30	13	46.4	1957	2 Q6Q1Y3	Q6Q1Y3 mus musculu
31	13	46.4	1957	2 Q63554	Q63554 rattus norv

32	13	46.4	1957	2 AAS45602	AAS45602 mus muscu
33	13	46.4	1958	2 P70276	P70276 mus musculu
34	13	46.4	1958	2 Q46669	Q46669 canis famli
35	12	42.9	1444	2 Q9UDM0	Q9UDM0 homo sapien
36	12	42.9	1791	2 Q8NDX3	Q8NDX3 homo sapien
37	12	42.9	1791	2 Q9UHE0	Q9UHE0 homo sapien
38	12	42.9	1791	2 Q9U133	Q9U133 homo sapien
39	11	39.3	1136	2 Q804F4	Q804F4 sternopygus
40	11	39.3	1717	2 Q90519	Q90519 fuqua rubrip
41	11	39.3	1949	2 Q9DRE3	Q9DRE3 brachydanio
42	11	39.3	1962	2 Q75RX9	Q75RX9 homo sapien
43	11	39.3	1962	2 BAD12085	BAD12085 homo sapi
44	11	39.3	1966	2 Q925G6	Q925G6 rattus norv
45	11	39.3	1976	2 Q63541	Q63541 rattus norv
46	11	39.3	1978	1 C1N8_MOUSE	C1N8_MOUSE
47	11	39.3	1978	1 Q884Z0	Q884Z0 rattus norv
48	11	39.3	1980	2 C1N8_HUMAN	C1N8_HUMAN
49	11	39.3	1988	2 Q884Z1	Q884Z1 rattus norv
50	11	39.3	2013	2 Q865W3	Q865W3 canis famli
51	11	39.3	2015	2 Q86UR3	Q86UR3 homo sapien
52	11	39.3	2015	2 Q81XC9	Q81XC9 homo sapien
53	11	39.3	2015	2 Q96J69	Q96J69 homo sapien
54	11	39.3	2016	1 C1N5_HUMAN	C1N5_HUMAN
55	11	39.3	2016	2 Q75RY0	Q75RY0 homo sapien
56	11	39.3	2016	2 BAD12084	BAD12084 homo sapi
57	11	39.3	2019	1 C1N5_RAT	C1N5_RAT
58	11	39.3	2019	2 Q9UJY9	Q9UJY9 mus musculu
59	11	39.3	2022	2 Q8WMP8	Q8WMP8 bos taurus
60	10	35.7	207	2 Q42421	Q42421 gallus gall
61	10	35.7	211	2 Q42418	Q42418 gallus gall
62	10	35.7	1784	2 Q25377	Q25377 lycoperico
63	9	32.1	286	2 Q00321	Q00321 lycoperico
64	9	32.1	286	2 Q40039	Q40039 hordeum vul
65	9	32.1	1765	2 Q88457	Q88457 rattus norv
66	9	32.1	1765	2 Q9JMD4	Q9JMD4 mus musculu
67	9	32.1	1765	2 Q9R053	Q9R053 mus musculu
68	8	28.6	60	2 Q8N0R3	Q8N0R3 pluteia xy
69	8	28.6	65	2 Q868A9	Q868A9 drosophila
70	8	28.6	80	2 Q7KQ27	Q7KQ27 heliothis v
71	8	28.6	269	2 Q18582	Q18582 caenorhabdi
72	8	28.6	305	2 Q7JPG9	Q7JPG9 drosophila
73	8	28.6	329	2 Q24719	Q24719 drosophila
74	8	28.6	362	2 Q7JN89	Q7JN89 drosophila
75	8	28.6	501	2 Q6DLT8	Q6DLT8 aedes albop
76	8	28.6	509	2 Q6DLT7	Q6DLT7 aedes albop
77	8	28.6	525	2 Q6DLT9	Q6DLT9 aedes albop
78	8	28.6	530	2 Q6DLU0	Q6DLU0 aedes aegypt
79	8	28.6	575	1 CYDD_BACSU	CYDD_BACSU
80	8	28.6	1087	2 Q9XYM6	Q9XYM6 lepidoptera
81	8	28.6	1538	2 Q7PE76	Q7PE76 drosophila
82	8	28.6	1618	2 Q8WMC7	Q8WMC7 anopheles g
83	8	28.6	1689	2 Q93135	Q93135 blattella g
84	8	28.6	1695	2 Q94584	Q94584 heliothis v
85	8	28.6	2031	2 Q01306	Q01306 blattella g
86	8	28.6	2031	2 Q01307	Q01307 blattella g
87	8	28.6	2051	2 Q86D17	Q86D17 pediculus h
88	8	28.6	2051	2 Q86D18	Q86D18 pediculus h
89	8	28.6	2051	2 Q86D19	Q86D19 pediculus h
90	8	28.6	2058	2 Q6DLT4	Q6DLT4 aedes albop
91	8	28.6	2064	2 Q6DLT3	Q6DLT3 aedes aegypt
92	8	28.6	2086	2 Q86M38	Q86M38 pediculus h
93	8	28.6	2104	2 Q25440	Q25440 musca domes
94	8	28.6	2105	2 Q25439	Q25439 musca domes
95	8	28.6	2108	2 Q94615	Q94615 musca domes
96	8	28.6	2131	1 C1N4_DROME	C1N4_DROME
97	8	28.6	2215	1 Q86D77	Q86D77 varroa deest
98	8	28.6	2223	2 Q7Q1V0	Q7Q1V0 anopheles g
99	8	28.6	2304	2 Q9W0Y8	Q9W0Y8 blattella g
100	8	28.6	2327	2 Q9W0Y8	Q9W0Y8 drosophila

ALIGNMENTS

```

RESULT 1
Q86X25 PRELIMINARY; PRT; 506 AA.
AC Q86X25;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DE Similar to sodium channel, voltage-gated, type II, alpha 2 polypeptide
   (Fragment).
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Brain;
RA Strausberg R.;
RL Submitted (FEB-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC047398; AAH47398.1; -.
DR HSP; P08104; I0G9.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0005261; P:cation channel activity; IEA.
DR GO; GO:0006812; P:cation transport; IEA.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR005821; Ion_trans.
DR Pfam; PF00520; M+channel_nlg.
DR Ion transport; Ion channel; Transmembrane; Transport.
KW NON_TER
SQ SEQUENCE 506 AA; 57147 MW; 39C32369D2A6D9 CRC64;

Query Match 100.0%; Score 28; DB 2; Length 506;
Best Local Similarity 100.0%; Pred. No. 2,5e-18;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 YMIFFVLVIFLGSPYLINILIAVVMAY 28
Db 401 YMIFFVLVIFLGSPYLINILIAVVMAY 428

RESULT 2
Q86X12 PRELIMINARY; PRT; 510 AA.
AC Q86X12;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Similar to sodium channel, voltage-gated, type II, alpha 2
   (Fragment).
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Brain;
RA Strausberg R.;
RL Submitted (MAR-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC048417; AAH48417.1; -.
DR HSP; P08104; I0G9.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0005261; P:cation channel activity; IEA.
DR GO; GO:0006812; P:cation transport; IEA.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR005821; Ion_trans.
DR Pfam; PF00520; M+channel_nlg.
DR Ion transport; Ion channel; Transmembrane; Transport.
KW NON_TER
SQ SEQUENCE 510 AA; 57660 MW; CFB153E259C32369 CRC64;

Query Match 100.0%; Score 28; DB 2; Length 510;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Best Local Similarity 100.0%; Pred. No. 2,6e-18;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 YMIFFVLVIFLGSPYLINILIAVVMAY 28
Db 401 YMIFFVLVIFLGSPYLINILIAVVMAY 428

RESULT 3
Q8IUU6 PRELIMINARY; PRT; 1981 AA.
AC Q8IUU6;
DT 01-MAR-2003 (TrEMBLrel. 23, Created)
DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE Voltage-gated sodium channel alpha 1 subunit.
GN Name=SCN1A;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Normal brain;
RA Ouchida M., Ohmori I.;
RL Submitted (DEC-2002) to the EMBL/GenBank/DBJ databases.
CC -1-SIMBLBLULAR LOCATION: Integral membrane protein (By similarity).
CC -1-SIMBLBLULAR LOCATION: Belongs to the sodium channel family.
DR EMBL; AB098335; BAC45228.1; -.
DR HSP; P04775; IBY5.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0001518; C:voltage-gated sodium channel complex; IEA.
DR GO; GO:0005261; P:cation channel activity; IEA.
DR GO; GO:0005248; P:voltage-gated sodium channel activity; IEA.
DR GO; GO:0006814; P:sodium ion transport; IEA.
DR GO; GO:0006812; P:sodium ion transport; IEA.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR002111; Cat_channel_Typl.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IQ_region.
DR InterPro; IPR005820; M+channel_nlg.
DR InterPro; IPR001696; Na_channel.
DR InterPro; IPR008051; Na_channel.
DR InterPro; IPR010525; Na_channel.
DR InterPro; IPR000100; Ribonuclease_P.
DR Pfam; PF00520; Ion_trans; 4.
DR Pfam; PF00612; IQ_1.
DR Pfam; PF06512; Na_trans_assoc; 1.
DR PRINTS; PR00170; NACHANNEL.
DR PRINTS; PR01664; NACHANNEL1.
DR SMART; SM00015; IQ_1.
DR PROSITE; PS00648; RIBONUCLEASE_P; UNKNOWN_1.
KW Ion transport; Ion channel; Sodium channel; Transmembrane;
   Transport; Voltage-gated channel.
SQ SEQUENCE 1981 AA; 226201 MW; BID6946D6491B7AD CRC64;

Query Match 100.0%; Score 28; DB 2; Length 1981;
Best Local Similarity 100.0%; Pred. No. 7,1e-18;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 YMIFFVLVIFLGSPYLINILIAVVMAY 28
Db 399 YMIFFVLVIFLGSPYLINILIAVVMAY 426

RESULT 4
CIN2 HUMAN STANDARD; PRT; 2005 AA.
ID CIN2 HUMAN
AC Q99250; Q14472; Q98ZC9; Q98ZD0;
DT 01-JUN-1994 (Rel. 29, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 05-JUL-2004 (Rel. 44, Last annotation update)
DE Sodium channel protein type II alpha subunit (Voltage-gated sodium

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DE channel alpha subunit Nav1.2) (Sodium channel protein, brain II alpha
DN subunit) (HSC II).
GN Name=SCN2A; Synonyms=SCN2A2, NAC2;
OS Homo sapiens (Human)
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A. (ISOFORM 1).
RC TISSUE=Brain;
RX MEDLINE=92390418; PubMed=1325650;
RA Ahmed C.M., Ware D.H., Lee S.C., Patten C.D., Ferrer-Montiel A.V.,
RA Schinder A.F., McPherson J.D., Wagner-McPherson C.B., Wasmuth J.O.,
RA Evans G.A., Montal M.;
RT "Primary structure, chromosomal localization, and functional
RT expression of a voltage-gated sodium channel from human brain.";
RT Proc. Natl. Acad. Sci. U.S.A. 89:8220-8224(1992).
RN [2]
RP SEQUENCE FROM N.A. (ISOFORMS 1 AND 2).
RX MEDLINE=21142400; PubMed=11245985; DOI=10.1016/S0378-1119(00)00594-1;
RA Kasai N., Fukushima K., Ueki Y., Prasad S., Nosakowski J.,
RA Sugata K.-I., Sugata A., Nishizaki K., Meyer N.C., Smith R.J.H.;
RT "Genomic structures of SCN2A and SCN3A - candidate genes for deafness
RT at the DFNA16 locus.";
RT Gene 264:113-122(2001).
RN [3]
RP SEQUENCE OF 1-89 FROM N.A.
RA Lu C.-M., Eichelberger J.S., Beckman M.L., Schade S.D., Brown G.B.;
RT "Isolation of the 5'-flanking region for human brain sodium channel
RT subtype II alpha-subunit (SCN2A).";
RT Submitted (APR-1998) to the EMBL/GenBank/DBJ databases.
RN [4]
RP SEQUENCE OF 1702-2005 FROM N.A.
RC TISSUE=Brain;
RX MEDLINE=92275082; PubMed=1317301;
RA Lu C.-M., Han J., Rado T.A., Brown G.B.;
RT "Differential expression of two sodium channel subtypes in human
RT brain.";
RT FEBS Lett. 303:53-58(1992).
RN [5]
RP SEQUENCE OF 1702-1772 FROM N.A.
RX MEDLINE=9110524; PubMed=184640;
RA Han J., Lu C.-M., Brown G.B., Rado T.A.;
RT "Direct amplification of a single dissected chromosomal segment by
RT polymerase chain reaction: a human brain sodium channel gene is on
RT chromosome 2q22-q23.";
RT Proc. Natl. Acad. Sci. U.S.A. 88:335-339(1991).
CC -1- FUNCTION: Mediates the voltage-dependent sodium ion permeability
CC of excitable membranes. Assuming opened or closed conformations in
CC response to the voltage difference across the membrane, the
CC protein forms a sodium-selective channel through which Na(+) ions
CC may pass in accordance with their electrochemical gradient.
CC -1- SUBUNIT: The sodium channel consists of a large polypeptide and 2-
CC 3 smaller ones. This sequence represents a large polypeptide.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
CC -1- ALTERNATIVE PRODUCTS:
CC Event=Alternative splicing; Named isoforms=2;
CC Name=1; Synonyms=Adult, 6A;
CC IsoId=G99250-1; Sequence=displayed;
CC Name=2; Synonyms=Neonatal, 6N;
CC IsoId=G99250-2; Sequence=VSP_001032;
CC -1- DOMAIN: The sequence contains 4 internal repeats, each with 5
CC hydrophobic segments (S1,S2,S3,S5,S6) and one positively charged
CC segment (S4). Segments S4 are probably the voltage-sensors and are
CC characterized by a series of positively charged amino acids at
CC every third position.
CC -1- SIMILARITY: Belongs to the sodium channel family.
CC -1- SIMILARITY: Contains 1 IQ domain.
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way

CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
CC or send an email to license@isb-sib.ch).
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FT REPEAT 1190 1504 III.
FT REPEAT 1513 1811 IV.
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FT TRANSMEM 157 176 S2 of repeat I.
FT TRANSMEM 190 208 S3 of repeat I.
FT TRANSMEM 215 234 S4 of repeat I.
FT TRANSMEM 251 274 S5 of repeat I.
FT TRANSMEM 402 427 S6 of repeat I.
FT TRANSMEM 754 778 S1 of repeat II.
FT TRANSMEM 790 813 S2 of repeat II.
FT TRANSMEM 822 841 S3 of repeat II.
FT TRANSMEM 848 867 S4 of repeat II.
FT TRANSMEM 884 904 S5 of repeat II.
FT TRANSMEM 958 983 S6 of repeat II.
FT TRANSMEM 1204 1237 S1 of repeat III.
FT TRANSMEM 1241 1266 S2 of repeat III.
FT TRANSMEM 1273 1294 S3 of repeat III.
FT TRANSMEM 1329 1350 S4 of repeat III.
FT TRANSMEM 1340 1367 S5 of repeat III.
FT TRANSMEM 1447 1473 S6 of repeat III.
FT TRANSMEM 1527 1550 S1 of repeat IV.
FT TRANSMEM 1562 1585 S2 of repeat IV.
FT TRANSMEM 1592 1615 S3 of repeat IV.
FT TRANSMEM 1626 1647 S4 of repeat IV.
FT TRANSMEM 1663 1685 S5 of repeat IV.
FT TRANSMEM 1752 1776 S6 of repeat IV.
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FT CARBOHYD 285 285 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 291 291 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 297 297 N-linked (GlcNAc...) (Potential).
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FT CARBOHYD 1368 1368 N-linked (GlcNAc...) (Potential).
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FT CONFLICT 524 524 /FTId=VSP_001032.
R -> L (in Ref. 1).

Query Match 100.0%; Score 28; DB 1; Length 2005;
Best Local Similarity 100.0%; Pred. No. 7, 2e-18;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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Db 401 YMFPLVIFLGSFYINILAVANAY 428

RESULT 5
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AC P04775;
DT 13-AUG-1987 (Rel. 05, Created)
DT 13-AUG-1987 (Rel. 05, Last sequence update)
DT 01-OCT-2004 (Rel. 45, Last annotation update)
DE Sodium channel protein type II alpha subunit (Voltage-gated sodium
DE channel alpha subunit Nav1.2) (Sodium channel protein, brain II alpha
DE subunit).
GN Name=Scn2a;
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxId=10116;

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RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=86146901; PubMed=3754035;
RA Noda M., Ikeda T., Kayano T., Suzuki H., Takeshima H., Kuraaki M.,
RA Takahashi H., Numa S.;
RT "Existence of distinct sodium channel messenger RNAs in rat brain.";
RL Nature 320:188-192(1986).
CC -1- FUNCTION: Mediates the voltage-dependent sodium ion permeability
CC of excitable membranes. Assuming opened or closed conformations in
CC response to the voltage difference across the membrane, the
CC protein forms a sodium-selective channel through which Na(+) ions
CC may pass in accordance with their electrochemical gradient.
CC -1- SUBUNIT: The sodium channel consists of a large polypeptide and 2-
CC 3 smaller ones. This sequence represents a large polypeptide.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
CC -1- DOMAIN: The sequence contains 4 internal repeats, each with 5
CC hydrophobic segments (S1,S2,S3,S5,S6) and one positively charged
CC segment (S4). Segments S4 are probably the voltage-sensors and are
CC characterized by a series of positively charged amino acids at
CC every third position.
CC -1- SIMILARITY: Belongs to the sodium channel family.
CC -1- SIMILARITY: Contains 1 IO domain.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL Outstation -
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CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC or send an email to license@sb-sib.ch).
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DR InterPro; IPR001682; Ca/Na_pore.
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DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IO_region.
DR InterPro; IPR005820; M_channel_nlg.
DR InterPro; IPR010526; Na_channel_assoc.
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DR PROSITE; PS50096; IO; 1.
KW 3D-structure; Glycoprotein; Ion transport; Ionic channel;
KW Multigene family; Repeat; Sodium channel; Transmembrane;
KW Voltage-gated channel.
KW
FT REPEAT 111 456 I.
FT REPEAT 741 1013 II.
FT REPEAT 1190 1504 III.
FT REPEAT 1513 1811 IV.
FT TRANSMEM 125 148 S1 of repeat I.
FT TRANSMEM 157 176 S2 of repeat I.
FT TRANSMEM 190 208 S3 of repeat I.
FT TRANSMEM 215 234 S4 of repeat I.
FT TRANSMEM 251 274 S5 of repeat I.
FT TRANSMEM 402 427 S6 of repeat I.
FT TRANSMEM 754 778 S1 of repeat II.
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FT TRANSMEM 822 841 S3 of repeat II.
FT TRANSMEM 848 867 S4 of repeat II.
FT TRANSMEM 884 904 S5 of repeat II.
FT TRANSMEM 958 983 S6 of repeat II.
FT TRANSMEM 1204 1237 S1 of repeat III.
FT TRANSMEM 1241 1266 S2 of repeat III.
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FT TRANSMEM 1299 1320 S4 of repeat III.
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FT TRANSMEM 1447 1473 S6 of repeat III.
FT TRANSMEM 1527 1550 S1 of repeat IV.
FT TRANSMEM 1562 1585 S2 of repeat IV.
FT TRANSMEM 1592 1615 S3 of repeat IV.

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FT TRANSMEM 1626 1647 S4 of repeat IV.
 FT TRANSMEM 1663 1685 S5 of repeat IV.
 FT TRANSMEM 1752 1776 S6 of repeat IV.
 FT DOMAIN 1905 1934 IO.
 FT CARBOHYD 212 212 N-linked (GlcNAc. . .) (Potential).
 FT CARBOHYD 285 285 N-linked (GlcNAc. . .) (Potential).
 FT CARBOHYD 291 291 N-linked (GlcNAc. . .) (Potential).
 FT CARBOHYD 297 297 N-linked (GlcNAc. . .) (Potential).
 FT CARBOHYD 303 303 N-linked (GlcNAc. . .) (Potential).
 FT CARBOHYD 308 308 N-linked (GlcNAc. . .) (Potential).
 FT CARBOHYD 340 340 N-linked (GlcNAc. . .) (Potential).
 FT CARBOHYD 604 604 N-linked (GlcNAc. . .) (Potential).
 FT CARBOHYD 624 624 N-linked (GlcNAc. . .) (Potential).
 FT CARBOHYD 883 883 N-linked (GlcNAc. . .) (Potential).
 FT CARBOHYD 1055 1055 N-linked (GlcNAc. . .) (Potential).
 FT CARBOHYD 1072 1072 N-linked (GlcNAc. . .) (Potential).
 FT CARBOHYD 1136 1136 N-linked (GlcNAc. . .) (Potential).
 FT CARBOHYD 1368 1368 N-linked (GlcNAc. . .) (Potential).
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OY 1 YMFPLVTFGSEFYINILIAVVMAY 28
 Db 401 YMFPLVTFGSEFYINILIAVVMAY 428

RESULT 6
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 AC P35498; Q16172; Q96LA3; Q9C008;
 DT 01-JUN-1994 (Rel. 29, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 05-JUL-2004 (Rel. 44, Last annotation update)
 DE Sodium channel protein type I alpha subunit (Voltage-gated sodium channel, alpha subunit Nav1.1) (Sodium channel protein, brain I alpha subunit).
 GN Name=SCN1A; Synonyms=SCN1, NAC1;
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 OC NCBI_taxid=9606;
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 RP SEQUENCE FROM N.A. (ISOFORM 1), AND VARIANTS GEFS+2 MET-875 AND HIS-1648.
 RX MEDLINE=20206553; PubMed=10742094;
 RA Escayg A., MacDonald B.T., Meisler M.H., Baulac S., Huberfeld G., An-Couffinkel I., Brice A., Leguern E., Mouldard B., Chaigne D., Buresi C., Malafosse A.;
 RT "Mutations of SCN1A, encoding a neuronal sodium channel, in two families with GEFS+2.";
 RT Nat. Genet. 24:343-345(2000).
 RU [2]
 RP SEQUENCE FROM N.A. (ISOFORM 2).
 RA Jeong S.-Y., Goto J., Kanazawa I.;
 RT "Cloning of cDNA for human voltage-gated sodium channel alpha subunit, SCN1A.";
 RU Submitted (JAN-2000) to the EMBL/GenBank/DBJ databases.
 RN [3]
 RP SEQUENCE FROM N.A. (ISOFORM 2).
 RC TISSUE=Brain;
 RA Sugawara T., Mazaki E.M., Yamakawa K.;
 RT "Homo sapiens neuronal voltage-gated sodium channel type I (Nav1.1) mRNA.";
 RU Submitted (JUL-2001) to the EMBL/GenBank/DBJ databases.
 RN [4]
 RP SEQUENCE FROM N.A. (ISOFORMS 1 AND 2), AND VARIANT THR-1067.
 RA Ouchida M., Omori I.;
 RT "Isoforms of human sodium channel SCN1A gene.";

RL Submitted (OCT-2002) to the EMBL/GenBank/DBJ databases.
 RN [5]
 RP SEQUENCE OF 1335-1428 FROM N.A.
 RX MEDLINE=94340991; PubMed=8062593;
 RA Malo M.S., Blanchard B.U., Andresen J.M., Srivastava K., Chen X.N., Li X., Jabe E.W., Korenberg J.R., Ingram V.M.;
 RT "Localization of a putative human brain sodium channel gene (SCN1A) to chromosome band 2q24.";
 RL Cytogenet. Cell Genet. 67:178-186(1994).
 RN [6]
 RP SEQUENCE OF 1518-1940 FROM N.A.
 RC TISSUE=Brain;
 RX MEDLINE=92275082; PubMed=1317301;
 RA Lu C.-M., Han J., Rado T.A., Brown G.B.;
 RT "Differential expression of two sodium channel subtypes in human brain.";
 RL FEBS Lett. 303:53-58(1992).
 RN [7]
 RP VARIANTS GEFS+2 VAL-188; LEU-1353 AND MET-1656.
 RX MEDLINE=21152274; PubMed=11254444;
 RA Wallace R.H., Scheffer I.E., Barnett S., Richards M., Dibbens L., Desai R.R., Lerman-Segie T., Lev D., Mazarib A., Brand N., Ben-Zeev B., Golkman I., Singh R., Kremidiotis G., Gardner A., Sutherland G.R., George A.L. Jr., Mulley J.C., Berkovic S.F.;
 RT "Neuronal sodium-channel alpha1-subunit mutations in generalized epilepsy with febrile seizures plus.";
 RL Am. J. Hum. Genet. 68:859-865(2001).
 RN [8]
 RP VARIANTS GEFS+2 ARG-1204.
 RX MEDLINE=21152275; PubMed=11254445;
 RA Escayg A., Heils A., MacDonald B.T., Haug K., Sander T., Meisler M.H.;
 RT "A novel SCN1A mutation associated with generalized epilepsy with febrile seizures plus -- and prevalence of variants in patients with epilepsy.";
 RL Am. J. Hum. Genet. 68:866-873(2001).
 RN [9]
 RP VARIANT SMEI PHE-986.
 RX MEDLINE=21257503; PubMed=11359211;
 RA Claes L., Del-Pavero J., Ceulemans B., Lagae L., Van Broeckhoven C., De Jonghe P.;
 RT "De novo mutations in the sodium-channel gene SCN1A cause severe myoclonic epilepsy of infancy.";
 RL Am. J. Hum. Genet. 68:1327-1332(2001).
 RN [10]
 RP VARIANTS GEFS+2 THR-1270.
 RX MEDLINE=21630138; PubMed=11756608;
 RA Abou-Khalil B., Ge Q., Desai R., Ryther R., Bazyk A., Bailey R., Haines J.L., Sutcliffe J.S., George A.L. Jr.;
 RT "Partial and generalized epilepsy with febrile seizures plus and a novel SCN1A mutation.";
 RL Neurology 57:2265-2272(2001).
 CC -1- FUNCTION: Mediates the voltage-dependent sodium ion permeability of excitable membranes. Assuming opened or closed conformations in response to the voltage difference across the membrane, the protein forms a sodium-selective channel through which Na(+) ions may pass in accordance with their electrochemical gradient.
 CC -1- SUBUNIT: The sodium channel consists of a large polypeptide and 2-3 smaller ones. This sequence represents a large polypeptide.
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
 CC -1- ALTERNATIVE PRODUCTS:
 CC Event=Alternative splicing; Named isoforms=2;
 CC Name=1;
 CC IsoId=P35498-1; Sequence=Displayed;
 CC Name=2;
 CC IsoId=P35498-2; Sequence=VSP_001031;
 CC Note=No experimental confirmation available;
 CC -1- DOMAIN: The sequence contains 4 internal repeats, each with 5 hydrophobic segments (51, 52, 53, 55, 56) and one positively charged segment (54). Segments 54 are probably the voltage-sensors and are characterized by a series of positively charged amino acids at every third position.
 CC -1- DISEASE: Defects in SCN1A are the cause of generalized epilepsy with febrile seizures plus type 2 (GEFS+2) [MIM:604233]. This

autosomal dominant disorder is characterized by febrile seizures in children and afebrile seizures in adults. Penetrance is incomplete and a large intrafamilial variability of the phenotype is observed.

-1- DISASE: Defects in SCN1A are a cause of severe myoclonic epilepsy in infancy (SMEI) [MIM:607208], a severe form of generalized epilepsy with febrile seizures. SMEI is a rare disorder characterized by normal development before onset, seizures beginning in the first year of life in the form of generalized or unilateral febrile clonic seizures, secondary appearance of myoclonic seizures, and occasionally partial seizures. It is associated with ataxia, slowed psychomotor development, and mental decline.

-1- SIMILARITY: Belongs to the sodium channel family.

-1- SIMILARITY: Contains 1 IQ domain.

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 DR EMBL; AY043484; AAK9360.1; -
 DR EMBL; AB093548; BAC21101.1; -
 DR EMBL; AB093549; BAC21102.1; -
 DR EMBL; S71446; AAB31605.1; -
 DR EMBL; X61863; CAA46439.1; -
 DR PIR; I52964; I52964. -
 DR PIR; S29184; S29184. -
 DR HSSP; P04775; IBY. -
 DR Genew; HGNC:10585; SCN1A. -
 DR MIM; 182389; -
 DR MIM; 604233; -
 DR MIM; 607208; -
 DR GO; GO:0016021; C:integral to membrane; NAS.
 DR GO; GO:0005248; P:voltage-gated sodium channel activity; NAS.
 DR GO; GO:0006814; P:sodium ion transport; NAS.
 DR InterPro; IPR001682; Ca/Na_pore.
 DR InterPro; IPR002111; Cat channel_Typl.
 DR InterPro; IPR005821; Ion trans.
 DR InterPro; IPR000048; IO region.
 DR InterPro; IPR005820; M+channel_nlg.
 DR InterPro; IPR001696; Na_channel.
 DR InterPro; IPR008051; Na_channel1.
 DR InterPro; IPR010526; Na_trans_assoc.
 DR Pfam; PF00520; Ion_trans_4.
 DR Pfam; PF00612; IQ_1.
 DR Pfam; PF06512; Na_trans_assoc; 1.
 DR PRINTS; PRO0170; NACHANNEL.
 DR PRINTS; PRO1664; NACHANNEL1.
 DR PROSITE; PS50096; IQ; FALSE_NEG.
 KW Alternative splicing; Disease mutation; Epilepsy; Glycoprotein;
 KW Ion transport; Ionic channel; Multigene family; Polymorphism; Repeat;
 KW Sodium channel; Transmembrane; Voltage-gated channel.
 FT REPEAT 110 454 I
 FT REPEAT 750 1022 II
 FT REPEAT 1200 1514 III
 FT REPEAT 1523 1821 IV
 FT TRANSMEM 14 147 S1 of repeat I (By similarity).
 FT TRANSMEM 156 175 S2 of repeat I (By similarity).
 FT TRANSMEM 189 207 S3 of repeat I (By similarity).
 FT TRANSMEM 214 233 S4 of repeat I (By similarity).
 FT TRANSMEM 250 273 S5 of repeat I (By similarity).
 FT TRANSMEM 400 425 S6 of repeat I (By similarity).
 FT TRANSMEM 763 787 S1 of repeat II (By similarity).
 FT TRANSMEM 799 822 S2 of repeat II (By similarity).
 FT TRANSMEM 831 850 S3 of repeat II (By similarity).
 FT TRANSMEM 857 876 S4 of repeat II (By similarity).
 FT TRANSMEM 893 913 S5 of repeat II (By similarity).

FT TRANSMEM 967 992 S6 of repeat II (By similarity).
 FT TRANSMEM 1214 1237 S1 of repeat III (By similarity).
 FT TRANSMEM 1251 1276 S2 of repeat III (By similarity).
 FT TRANSMEM 1283 1304 S3 of repeat III (By similarity).
 FT TRANSMEM 1309 1330 S4 of repeat III (By similarity).
 FT TRANSMEM 1350 1377 S5 of repeat III (By similarity).
 FT TRANSMEM 1457 1483 S6 of repeat III (By similarity).
 FT TRANSMEM 1537 1560 S1 of repeat IV (By similarity).
 FT TRANSMEM 1572 1595 S2 of repeat IV (By similarity).
 FT TRANSMEM 1602 1625 S3 of repeat IV (By similarity).
 FT TRANSMEM 1636 1657 S4 of repeat IV (By similarity).
 FT TRANSMEM 1673 1695 S5 of repeat IV (By similarity).
 FT TRANSMEM 1762 1786 S6 of repeat IV (By similarity).
 FT CARBOHYD 211 211 N-linked (GlcNAc...) (Potential).

Query Match 100.0%; Score 28; DB 1; Length 2009;
 Best local similarity 100.0%; Pred. No. 7,2e-18;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 YWIFVVLVIFGSEFYLLNLIIAVVAMAY 28
 DB 399 YWIFVVLVIFGSEFYLLNLIIAVVAMAY 426

RESULT 7
 ID CINI_RAT STANDARD; PRT; 2009 AA.
 AC P04774;
 DT 13-AUG-1987 (Rel. 05, Created)
 DT 13-AUG-1987 (Rel. 05, Last sequence update)
 DT 05-JUL-2004 (Rel. 44, Last annotation update)
 DE Sodium channel protein type I alpha subunit (voltage-gated sodium channel, alpha subunit Nav1.1) (Sodium channel protein, brain I alpha subunit).
 DE Name=Scn1a;
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sclurognathu; Muridae; Murinae; Rattus.
 OX NCBI_TaxID=10116;
 [1]
 RN Nucleo 320:188-192(1986).
 RP SEQUENCE FROM N.A.
 RX MEDLINE=86146901; PubMed=3754035;
 RA Noda M., Ikeda T., Kayano T., Suzuki H., Takeshima H., Kurasaki M., Takahashi H., Numa S.;
 RT "Existence of distinct sodium channel messenger RNAs in rat brain.";
 RL Nature 320:188-192(1986).
 [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=87311395; PubMed=2442385;
 RA Noda M., Numa S.;
 RT "Structure and function of sodium channel.";
 RL J. Recept. Res. 7:467-497(1987).
 CC -1- FUNCTION: Mediates the voltage-dependent sodium ion permeability of excitable membranes. Assuming opened or closed conformations in response to the voltage difference across the membrane, the protein forms a sodium-selective channel through which Na(+) ions may pass in accordance with their electrochemical gradient.
 CC -1- SUBUNIT: The sodium channel consists of a large polypeptide and 2-3 smaller ones. This sequence represents a large polypeptide.
 CC -1- DOMAIN: The sequence contains 4 internal repeats, each with 5 hydrophobic segments (S1,S2,S3,S5,S6) and one positively charged segment (S4). Segments S4 are probably the voltage-sensors and are characterized by a series of positively charged amino acids at every third position.
 CC -1- SIMILARITY: Belongs to the sodium channel family.
 CC -1- SIMILARITY: Contains 1 IQ domain.

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CC EMBL, X03638; CAA37286.1; -
DR EMBL, M22253; AAA79965.1; -
DR PIR, A25019; A25019.
DR HSSP, P04775; 1BYV.
DR RGD, 69364; Scnla.
DR InterPro; IPR001682; Ca/Na.pore.
DR InterPro; IPR002111; Cat_channel_Tryp.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IQ_region.
DR InterPro; IPR005820; M+channel_nlg.
DR InterPro; IPR001696; Na_channel.
DR InterPro; IPR008051; Na_channel.
DR InterPro; IPR010526; Na_trans_assoc.
DR Pfam; PF005120; Ion_trans_4.
DR Pfam; PF00612; IQ_1.
DR Pfam; PF06512; Na_trans_assoc; 1.
DR PRINTS; PR00170; NACHANNEL.
DR PRINTS; PR01664; NACHANNEL.
DR PROSITE; PS50096; IQ_FALSE_NEG.
DR Glycoprotein; Ion transport; Ionic channel; Multigene family; Repeat;
KW Sodium channel; Transmembrane; Voltage-gated channel.
FT REPEAT 110 454
FT REPEAT 1200 1032
FT REPEAT 1200 1514
FT REPEAT 1523 1821
FT TRANSMEM 124 147
FT TRANSMEM 156 175
FT TRANSMEM 189 207
FT TRANSMEM 214 233
FT TRANSMEM 250 273
FT TRANSMEM 400 425
FT TRANSMEM 763 787
FT TRANSMEM 799 822
FT TRANSMEM 831 850
FT TRANSMEM 857 876
FT TRANSMEM 883 913
FT TRANSMEM 967 992
FT TRANSMEM 1214 1237
FT TRANSMEM 1251 1276
FT TRANSMEM 1283 1304
FT TRANSMEM 1309 1330
FT TRANSMEM 1350 1377
FT TRANSMEM 1457 1483
FT TRANSMEM 1537 1560
FT TRANSMEM 1572 1595
FT TRANSMEM 1602 1625
FT TRANSMEM 1636 1657
FT TRANSMEM 1673 1695
FT TRANSMEM 1762 1786
FT CARBOHYD 211 211
FT CARBOHYD 284 284
FT CARBOHYD 295 295
FT CARBOHYD 301 301
FT CARBOHYD 306 306
FT CARBOHYD 338 338
FT CARBOHYD 601 601
FT CARBOHYD 621 621
FT CARBOHYD 681 681
FT CARBOHYD 892 892
FT CARBOHYD 1060 1060
FT CARBOHYD 1064 1064
FT CARBOHYD 1080 1080
FT CARBOHYD 1146 1146
FT CARBOHYD 1378 1378
FT CARBOHYD 1392 1392
FT CARBOHYD 1403 1403
SQ SEQUENCE 2009 AA; 228769 MW; 6808466F636837B CRC64;

Query Match 100.0%; Score 28; DB 1; Length 2009;
Best Local Similarity 100.0%; Pred. No. 7.2e-18;

Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 YMIFFVLVIFLGSPYLINILAVVMAY 28
DB 399 YMIFFVLVIFLGSPYLINILAVVMAY 426

RESULT 8
ID 042419 PRELIMINARY; PRT; 213 AA.
AC 042419; 01-JAN-1998 (TREMBlrel. 05, Created)
DT 01-JAN-1998 (TREMBlrel. 05, Last sequence update)
DE 01-JUN-2003 (TREMBlrel. 24, Last annotation update)
DE Voltage-gated sodium channel I (Fragment).
OS Gallus gallus (Chicken).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Archosauria; Aves; Neognathae; Galliformes; Phasianidae; Phasianinae;
OC Gallus.
OX NCBI_TaxID=9031;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Dorsal root ganglion;
RX MEDLINE=98024190; PubMed=9356508;
RA Friedel R.H., Schnuerch H., Stubbusch J., Barde Y.A.;
RT "Identification of genes differentially expressed by nerve growth
factor- and neurotrophin-3-dependent sensory neurons";
RL Proc. Natl. Acad. Sci. U.S.A. 94:12670-12675(1997).
DR EMBL; AJ001489; CAA04784.1; -
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0005261; P:cation channel activity; IEA.
DR GO; GO:006812; P:cation transport; IEA.
DR InterPro; IPR001682; Ca/Na.pore.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR005820; M+channel_nlg.
DR Pfam; PF00520; Ion_trans_1.
KW Ion transport; Ionic channel; Transmembrane; Transport.
FT NON_TER 1 213
FT NON_TER 1 213
SQ SEQUENCE 213 AA; 24397 MW; DAF3B6AE4E8B47ED CRC64;

Query Match 75.0%; Score 21; DB 2; Length 213;
Best Local Similarity 100.0%; Pred. No. 5.7e-12;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 YMIFFVLVIFLGSPYLINIL 21
DB 193 YMIFFVLVIFLGSPYLINIL 213
RESULT 9
ID 042420 PRELIMINARY; PRT; 225 AA.
AC 042420; 01-JAN-1998 (TREMBlrel. 05, Created)
DT 01-JAN-1998 (TREMBlrel. 05, Last sequence update)
DT 01-JUN-2003 (TREMBlrel. 24, Last annotation update)
DE Voltage-gated sodium channel II (Fragment).
OS Gallus gallus (Chicken).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Archosauria; Aves; Neognathae; Galliformes; Phasianidae; Phasianinae;
OC Gallus.
OX NCBI_TaxID=9031;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Dorsal root ganglion;
RX MEDLINE=98024190; PubMed=9356508;
RA Friedel R.H., Schnuerch H., Stubbusch J., Barde Y.A.;
RT "Identification of genes differentially expressed by nerve growth
factor- and neurotrophin-3-dependent sensory neurons";
RL Proc. Natl. Acad. Sci. U.S.A. 94:12670-12675(1997).
DR EMBL; AJ001490; CAA04785.1; -
DR GO; GO:0016021; C:integral to membrane; IEA.

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DR GO: GO:0005261; P: cation channel activity; IEA.
DR GO: GO:0006812; P: cation transport; IEA.
DR InterPro: IPR001682; Ca/Na pore.
DR InterPro: IPR005821; Ion trans.
DR InterPro: IPR005820; M+channel_nlg.
DR Pfam: PF00520; Ion trans; 1.
KW Ion transport; Ionic channel; Transmembrane; Transport.
FT NON TER 1 1
FT 225 225
SQ SEQUENCE 225 AA; 25531 MW; 65B122E582F3023E CRC64;

Query Match
Best Local Similarity 75.0%; Score 21; DB 2; Length 225;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 YMFPLVFLGSPYLNLIL 21
DB 205 YMFPLVFLGSPYLNLIL 225

RESULT 10
Q9IBD6 PRELIMINARY; PRT; 247 AA.
AC Q9IBD6;
DT 01-OCT-2000 (TREMBlrel. 15, Created)
DT 01-OCT-2000 (TREMBlrel. 15, Last sequence update)
DT 01-JUN-2003 (TREMBlrel. 24, Last annotation update)
DE Skeletal muscle voltage-gated sodium channel (Fmna2) (Fragment).
OS Takifugu pardalis (Pufferfish).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
OC Acanthomorphi; Acanthopterygii; Percomorphi; Tetraodontiformes;
OC Tetraodontidae; Tetraodontidae; Takifugu.
RX NCBI_TaxID=98921;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Skeletal muscle;
RK MEDLINE=20090650; PubMed=10623632;
RA Yocsu-Yamashita M., Nishimori K., Mitani Y., Isejima M., Sugimoto A.,
RA Yano M. T.;
RT "Binding properties of 3H-PbTx-3 and 3H-saxitoxin to brain membranes
RT and to skeletal muscle membranes of puffer fish Takifugu pardalis, and the
RT primary structure of a voltage-gated Na+ channel alpha-subunit (Fmna1)
RT from skeletal muscle of F. pardalis."
RL Biochem. Biophys. Res. Commun. 267:403-412 (2000).
DR EMBL: AB032022; BAA90308.1; -.
DR GO: GO:0016021; C: integral to membrane; IEA.
DR GO: GO:0005261; P: cation channel activity; IEA.
DR GO: GO:0006812; P: cation transport; IEA.
DR InterPro: IPR001682; Ca/Na pore.
DR InterPro: IPR005821; Ion trans.
DR InterPro: IPR005820; M+channel_nlg.
DR Pfam: PF00520; Ion trans; 1.
KW Ion transport; Ionic channel; Transmembrane; Transport.
FT NON TER 1 1
FT 247 247
SQ SEQUENCE 247 AA; 27998 MW; 08B8C1CECEFE05E1F5 CRC64;

Query Match
Best Local Similarity 75.0%; Score 21; DB 2; Length 247;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 VFELGSPYLNLILVVMAY 28
DB 223 VFELGSPYLNLILVVMAY 243

RESULT 11
CINA_ELEBL STANDARD; PRT; 1820 AA.
AC P02719;
DT 21-JUL-1986 (Rel. 01, Created)
DT 21-JUL-1986 (Rel. 01, Last sequence update)

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DT 29-MAR-2004 (Rel. 43, Last annotation update)
DE Sodium channel protein (Na+ channel)
OS Electrophorus electricus (Electric eel)
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Gymnotiformes;
OC Electrophoridae; Electrophorus.
RX NCBI_TaxID=8005;
RN [1]
RP SEQUENCE FROM N.A.
RC MEDLINE=85061498; PubMed=6209577;
RA Noda M., Shimizu S., Tanabe T., Takai T., Kayano T., Ikeda T.,
RA Takahashi H., Nakayama H., Kanaoka Y., Minamino N., Kangawa K.,
RA Matsuo H., Ratterey M.A., Hirose T., Inayama S., Hayashida H.,
RA Miyata T., Numa S.;
RT "Primary structure of Electrophorus electricus sodium channel deduced
RT from cDNA sequence."
RL Nature 312:121-127 (1984).
RN [2]
RP SEQUENCE FROM N.A.
RC MEDLINE=87311395; PubMed=2442385;
RA Noda M., Numa S.;
RT "Structure and function of sodium channel."
RL J. Recept. Res. 7:467-497 (1987).
CC -1- FUNCTION: Mediates the voltage-dependent sodium ion permeability
CC of excitable membranes. Assuming opened or closed conformations in
CC response to the voltage difference across the membrane, the
CC protein forms a sodium-selective channel through which Na(+) ions
CC may pass in accordance with their electrochemical gradient.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
CC -1- DOMAIN: The sequence contains 4 internal repeats, each with 5
CC hydrophobic segments (S1, S2, S3, S5, S6) and one positively charged
CC segment (S4). Segments S4 are probably the voltage-sensors and are
CC characterized by a series of positively charged amino acids at
CC every third position.
CC -1- MISCELLANEOUS: Available data suggest that activation and
CC inactivation gates are located near the cytoplasmic surface of the
CC membrane. It is hypothesized that residues 802-806, 847-857, 894-
CC 910, and 942-955 might, in conjunction with the positively charged
CC residues of S4, act as a voltage sensor involved with the
CC activation gate.
CC -1- SIMILARITY: Belongs to the sodium channel family.
CC -1- SIMILARITY: Contains 1 IQ domain.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL: X01119; CAA25587.1; -.
DR EMBL: M22252; AAA79960.1; -.
DR PIR: A03178; CHER.
DR HSP: P04775; IBY.
DR InterPro: IPR001682; Ca/Na pore.
DR InterPro: IPR002111; Cat channel_TripL.
DR InterPro: IPR005821; Ion trans.
DR InterPro: IPR000048; IQ region.
DR InterPro: IPR005820; M+channel_nlg.
DR InterPro: IPR001682; Na channel.
DR InterPro: IPR00526; Na_trans.
DR Pfam: PF00520; Ion trans; 4.
DR Pfam: PF00612; IQ; 1.
DR Pfam: PF06512; Na_trans_assoc; 1.
DR PRINTS: PR00170; NACHANNEL.
DR PROSITE: PS50096; IQ; FALSE_NEG.
DR Glycoprotein; Ion transport; Ionic channel; Repeat; Sodium channel;
KW Transmembrane; Voltage-gated channel.
FT REPEAT 111 419
FT REPEAT 111 807
FT REPEAT 989 1281
FT REPEAT 1311 1587
FT

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FT TRANSMEM 118 138 S1 of repeat I.
FT TRANSMEM 150 171 S2 of repeat I.
FT TRANSMEM 177 197 S3 of repeat I.
FT TRANSMEM 204 224 S4 of repeat I.
FT TRANSMEM 244 264 S5 of repeat I.
FT TRANSMEM 285 342 Non-homologous region of repeat I.
FT TRANSMEM 379 402 S6 of repeat I.
FT TRANSMEM 428 558 S1 of repeat II.
FT TRANSMEM 600 620 S2 of repeat II.
FT TRANSMEM 626 643 S3 of repeat II.
FT TRANSMEM 651 671 S4 of repeat II.
FT TRANSMEM 691 711 S5 of repeat II.
FT TRANSMEM 767 790 S6 of repeat II.
FT TRANSMEM 1005 1025 S1 of repeat III.
FT TRANSMEM 1038 1058 S2 of repeat III.
FT TRANSMEM 1066 1086 S3 of repeat III.
FT TRANSMEM 1092 1112 S4 of repeat III.
FT TRANSMEM 1132 1152 S5 of repeat III.
FT TRANSMEM 1172 1194 Non-homologous region of repeat III.
FT TRANSMEM 1244 1264 S6 of repeat III.
FT TRANSMEM 1321 1341 S1 of repeat IV.
FT TRANSMEM 1353 1376 S2 of repeat IV.
FT TRANSMEM 1381 1398 S3 of repeat IV.
FT TRANSMEM 1417 1437 S4 of repeat IV.
FT TRANSMEM 1454 1474 S5 of repeat IV.
FT TRANSMEM 1490 1505 Non-homologous region of repeat IV.
FT TRANSMEM 1544 1567 S6 of repeat IV.
FT CAROXYD 278 278 N-linked (GlcNAc. . .) (Potential).
FT CAROXYD 288 288 N-linked (GlcNAc. . .) (Potential).
FT CAROXYD 317 317 N-linked (GlcNAc. . .) (Potential).
FT CAROXYD 591 591 N-linked (GlcNAc. . .) (Potential).
FT CAROXYD 690 690 N-linked (GlcNAc. . .) (Potential).
FT CAROXYD 797 797 N-linked (GlcNAc. . .) (Potential).
FT CAROXYD 1160 1160 N-linked (GlcNAc. . .) (Potential).
FT CAROXYD 1174 1174 N-linked (GlcNAc. . .) (Potential).
SQ SEQUENCE 1820 AA; 208331 MW; 1B271F626E057864 CRC64;

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Query Match 75.0%; Score 21; DB 1; Length 1820;
 Best Local Similarity 100.0%; Pred. No. 2.9e-11;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 8 VIFLGSFYINILAVVAVAY 28
 |||||
 Db 385 VIFLGSFYINILAVVAVAY 405

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RESULT 12
Q9IBF1 PRELIMINARY; PRT; 1880 AA.
ID Q9IBF1
AC Q9IBF1:
DT 01-OCT-2000 (TrEMBLrel. 15, Created)
DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE Voltage-gated sodium channel.
OS Takifugu pardalis (Pufferfish).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Euteleostei;
OC Acanthomorpha; Acanthopterygii; Percomorph; Tetraodontiformes;
OC Tetraodontidae; Tetraodontidae; Takifugu.
OX NCBI_TaxID=98921;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=skelatal muscle;
RX MEDLINE=20090650; PubMed=10623632;
RA Yotsubo-Yamashita M., Nishimori K., Nitanai Y., Iseura M., Sugimoto A.,
RA Yasumoto T.;
RT "Binding properties of 3H-PbTx-3 and 3H-saxitoxin to brain membranes
RT and to skeletal muscle membranes of puffer fish Takifugu pardalis, and the
RT primary structure of a voltage-gated Na+ channel alpha-subunit (fMaai)
RT from skeletal muscle of F. pardalis.";
RL Biochem. Biophys. Res. Commun. 267:403-412(2000).
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
CC -1- SIMILARITY: Belongs to the sodium channel family.

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DR EMBL; AB030482; BAA90398.1; -.
DR HSSP; P04775; 1BYX.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0001518; C:voltage-gated sodium channel complex; IEA.
DR GO; GO:0005261; P:cation channel activity; IEA.
DR GO; GO:0005248; P:voltage-gated sodium channel activity; IEA.
DR GO; GO:0006812; P:cation transport; IEA.
DR GO; GO:0006814; P:sodium ion transport; IEA.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR002111; Cat_channel_Trtpl.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IQ_region.
DR InterPro; IPR005820; M+channel_nlg.
DR InterPro; IPR001696; Na_channel1.
DR InterPro; IPR010526; Na_trans_assoc.
DR Pfam; PF00520; Ion_trans_4.
DR Pfam; PF0612; IQ_1.
DR Pfam; PF06512; Na_trans_assoc; 1.
DR PRINTS; PR00170; NACHANNEL.
DR PROSITE; PS50096; IQ_1.
KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
KW Transport; Voltage-gated channel.
SQ SEQUENCE 1880 AA; 212084 MW; 406483C6C3D43E02 CRC64;

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Query Match 75.0%; Score 21; DB 2; Length 1880;
 Best Local Similarity 100.0%; Pred. No. 2.9e-11;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 8 VIFLGSFYINILAVVAVAY 28
 |||||
 Db 405 VIFLGSFYINILAVVAVAY 425

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RESULT 13
Q1S858 PRELIMINARY; PRT; 1977 AA.
ID Q1S858
AC Q1S858;
DT 01-NOV-1996 (TrEMBLrel. 01, Created)
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE Sodium channel alpha subunit.
GN Name=hmr-Na;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Thyroid;
RX MEDLINE=95237189; PubMed=7720699;
RA Klugbauer N., Lacinova L., Flockeiz V., Hofmann F.;
RT "Structure and functional expression of a new member of the
RT tetrodotoxin-sensitive voltage-activated sodium channel family from
RT human neuroendocrine cells.";
RL EMBO J. 14:1084-1090(1995).
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
CC -1- SIMILARITY: Belongs to the sodium channel family.
DR EMBL; X82835; CAAS8042.1; -.
DR PIR; S54771; S54771.
DR HSSP; P04775; 1BYX.
DR Genew; HGNC:10597; SCN9A.
DR GO; GO:0005248; P:voltage-gated sodium channel activity; TAS.
DR GO; GO:0006814; P:sodium ion transport; TAS.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR002111; Cat_channel_Trtpl.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IQ_region.
DR InterPro; IPR005820; M+channel_nlg.
DR InterPro; IPR001696; Na_channel1.
DR InterPro; IPR010526; Na_trans_assoc.
DR Pfam; PF00520; Ion_trans_4.
DR Pfam; PF0612; IQ_1.
DR Pfam; PF06512; Na_trans_assoc; 1.

```

DR PRINTS; PRO0170; NACHANNEL.
 DR SMART; SM00015; IQ; 1.
 KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
 SQ SEQUENCE 1977 AA; 225195 MW; 17D67CBG32BC15FB CRC64;

Query Match 75.0%; Score 21; DB 2; Length 1977;
 Best Local Similarity 100.0%; Pred. No. 3.1e-11;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 VIFLGSFYINLILAVVAMAY 28
 DB 385 VIFLGSFYINLILAVVAMAY 405

RESULT 14
 Q28644
 ID Q28644 PRELIMINARY; PRT; 1984 AA.
 AC Q28644;
 DT 01-NOV-1996 (TREMBlrel. 01, Created)
 DT 01-NOV-1996 (TREMBlrel. 01, Last sequence update)
 DT 01-MAR-2004 (TREMBlrel. 26, Last annotation update)
 DE Sodium channel alpha-subunit.
 OS Oryzocolagus cuniculus (Rabbit).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Lagomorpha; Leporidae; Oryctolagus.
 OX NCBI_TaxID=9986;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=New Zealand White; TISSUE=Sciatic nerve;
 RX MEDLINE=96074641; PubMed=7479931;
 RA Belcher S.M., Zerial C.A., Levenson R., Ritchie J.M., Howe J.R.;
 RT "Cloning of a sodium channel alpha subunit from rabbit Schwann
 cells.";
 RL Proc. Natl. Acad. Sci. U.S.A. 92:11034-11038(1995).
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
 CC -1- SIMILARITY: Belongs to the sodium channel family.
 DR EMBL; U35238; AAA89159.1; -.
 DR HSSP; P04775; 1BYX.
 DR GO; GO:0016021; C:Integral to membrane; IEA.
 DR GO; GO:0001518; C:voltage-gated sodium channel complex; IEA.
 DR GO; GO:0005261; F:cation channel activity; IEA.
 DR GO; GO:0005248; F:voltage-gated sodium channel activity; IEA.
 DR GO; GO:0006812; P:cation transport; IEA.
 DR GO; GO:0006814; P:sodium ion transport; IEA.
 DR InterPro; IPR001682; Ca/Na_pore.
 DR InterPro; IPR002111; Cat_channel_TrpL.
 DR InterPro; IPR005821; Ion_trans.
 DR InterPro; IPR000048; IQ_region.
 DR InterPro; IPR005820; M+channel_nlg.
 DR InterPro; IPR001696; Na_channel.
 DR InterPro; IPR010526; Na_trans_assoc.
 DR Pfam; PF00520; Ion_trans_4.
 DR Pfam; PF00612; IQ_1.
 DR Pfam; PF06512; Na_trans_assoc; 1.
 DR PRINTS; PRO0170; NACHANNEL.
 DR SMART; SM00015; IQ; 1.
 KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
 KW Transport; Voltage-gated channel.
 SQ SEQUENCE 1984 AA; 225748 MW; 98F76860C9866AA0 CRC64;

Query Match 75.0%; Score 21; DB 2; Length 1984;
 Best Local Similarity 100.0%; Pred. No. 3.1e-11;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 VIFLGSFYINLILAVVAMAY 28
 DB 383 VIFLGSFYINLILAVVAMAY 403

RESULT 15
 008562
 ID 008562 PRELIMINARY; PRT; 1984 AA.

AC 008562;
 DT 01-JUL-1997 (TREMBlrel. 04, Created)
 DT 01-JUL-1997 (TREMBlrel. 04, Last sequence update)
 DT 05-JUL-2004 (TREMBlrel. 27, Last annotation update)
 DE PNI (Voltage-gated sodium channel) (Fragment).
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 OX NCBI_TaxID=10116;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC MEDLINE=97188502; PubMed=9037087;
 RA Toledo-Aral J.J., Moss B.L., He Z.J., Kozowski A.G., Whisenand T.,
 RA Levinson S.R., Wolf J.J., Silos-Santiago I., Halsegoua S., Mandel G.;
 RT "Identification of PNI, a predominant voltage-dependent sodium channel
 expressed principally in peripheral neurons.";
 RL Proc. Natl. Acad. Sci. U.S.A. 94:1527-1532(1997).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC MEDLINE=97007982; PubMed=8854872;
 RA Kozak C.A., Sangameswaran L.;
 RT "Genetic mapping of the peripheral sodium channel genes, Scn9a and
 Scn10a, in the mouse.";
 RL Mamm. Genome 7:787-788(1996).
 RN [3]

RP SEQUENCE FROM N.A.
 RA Sangameswaran L., Fish L.M., Koch B.D., Rabert D.K., Delgado S.G.,
 RA Ilinicka M., Jakeman L.B., Novakovic S., Wong K., Sze P., Tzounakia E.,
 RA Stewart G.R., Herman R.C., Chan H., Eglén R.M., Hunter J.C.;
 RT "A novel tetrodotoxin-sensitive, voltage-gated sodium channel
 expressed in rat and human dorsal root ganglia.";
 RL J. Biol. Chem. 0:0-0(1997).
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
 CC -1- SIMILARITY: Belongs to the sodium channel family.
 DR EMBL; U79568; AAB50403.1; -.
 DR EMBL; AF000368; AAB80701.1; -.
 DR HSSP; P04775; 1BYX.
 DR GO; GO:0016021; C:Integral to membrane; IEA.
 DR GO; GO:0001518; C:voltage-gated sodium channel complex; IEA.
 DR GO; GO:0005261; F:cation channel activity; IEA.
 DR GO; GO:0005248; F:voltage-gated sodium channel activity; IEA.
 DR GO; GO:0006812; P:cation transport; IEA.
 DR GO; GO:0006814; P:sodium ion transport; IEA.
 DR InterPro; IPR001682; Ca/Na_pore.
 DR InterPro; IPR002111; Cat_channel_TrpL.
 DR InterPro; IPR005821; Ion_trans.
 DR InterPro; IPR000048; IQ_region.
 DR InterPro; IPR005820; M+channel_nlg.
 DR InterPro; IPR001696; Na_channel.
 DR InterPro; IPR010526; Na_trans_assoc.
 DR Pfam; PF00520; Ion_trans_4.
 DR Pfam; PF00612; IQ_1.
 DR Pfam; PF06512; Na_trans_assoc; 1.
 DR PRINTS; PRO0170; NACHANNEL.
 DR SMART; SM00015; IQ; 1.
 KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
 KW Transport; Voltage-gated channel.
 FT NON_TER 1984
 SQ SEQUENCE 1984 AA; 226037 MW; 386C38B9B5097091 CRC64;

Query Match 75.0%; Score 21; DB 2; Length 1984;
 Best Local Similarity 100.0%; Pred. No. 3.1e-11;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 VIFLGSFYINLILAVVAMAY 28
 DB 385 VIFLGSFYINLILAVVAMAY 405

RESULT 16
 025150
 ID 025150 PRELIMINARY; PRT; 2049 AA.

DT 01-NOV-1996 (TRENBLREL. 01, Created)
 DT 01-NOV-1996 (TRENBLREL. 01, Last sequence update)
 DT 01-MAR-2004 (TRENBLREL. 26, Last annotation update)
 DE Voltage-gated sodium channel.
 GN Name=tuna1;
 OS Halocynthia roretzi (Sea squirt).
 OC Eukaryota; Metazoa; Chordata; Urochordata; Ascidiacea;
 OC Stolidobranchia; Pyrosidae; Halocynthia.
 OX NCBI_TaxID=7729;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=95033215; PubMed=7946338;
 RA Okamura Y., Ono F., Okagaki R., Chong J., Mandel G.;
 RT "Neural expression of a sodium channel gene requires cell-specific
 interactions";
 RL Neuron 13:937-948(1994).
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
 CC -1- SIMILARITY: Belongs to the sodium channel family.
 DR EMBL, D17311; BAA04133.1; -.
 DR PIR, T43161; T43161.
 DR HSSP; P04775; 1BYV.
 DR GO; GO:0016021; C:integral to membrane; IEA.
 DR GO; GO:0001518; C:voltage-gated sodium channel complex; IEA.
 DR GO; GO:0005261; F:cation channel activity; IEA.
 DR GO; GO:0005248; F:voltage-gated sodium channel activity; IEA.
 DR GO; GO:0006812; P:cation transport; IEA.
 DR GO; GO:0006814; P:sodium ion transport; IEA.
 DR InterPro; IPR001682; Ca/Na_Pore.
 DR InterPro; IPR002111; Cat_channel_TpL.
 DR InterPro; IPR005821; Ion_trans.
 DR InterPro; IPR000048; IQ_region.
 DR InterPro; IPR005820; M_channel_nlg.
 DR InterPro; IPR001696; Na_channel_nlg.
 DR Pfam; PF00520; Ion_trans_4.
 DR Pfam; PF00612; IQ_1.
 DR Pfam; PF06512; Na_trans_assoc; 1.
 DR PRINTS; PR00170; NACHANNEL.
 DR SMART; SM00015; IQ_1.
 KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
 KW Transport; Voltage-gated channel.
 SQ SEQUENCE 2049 AA; 233443 MW; F7DB3578105B73FB CRC64;
 Query Match 75.0%; Score 21; DB 2; Length 2049;
 Best Local Similarity 100.0%; Fred.No. 3.1e-11;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 8 VIFLGSFYILNLIILAVVMAY 28
 |||||
 DB 449 VIFLGSFYILNLIILAVVMAY 469

RESULT 17
 CIN4_HUMAN STANDARD; PRT; 1836 AA.
 ID CIN4_HUMAN 015478; Q16447; Q726B1;
 AC P35459; 015478; 015478; 015478;
 DT 01-JUN-1994 (Rel. 29, Created)
 DT 29-MAR-2004 (Rel. 43, Last sequence update)
 DT 05-JUL-2004 (Rel. 44, Last annotation update)
 DE Sodium channel protein type IV alpha subunit (Voltage-gated sodium
 DE channel alpha subunit Nav1.4) (Sodium channel protein, skeletal muscle
 DE alpha-subunit).
 GN Name=SCN4A;
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A., AND VARIANT ASP-559.
 RX TISSUE=Skeletal muscle;
 RC MEDLINE=92246457; PubMed=1315496;
 RA George A.L. Jr., Komisarof J., Kallen R.G., Barchi R.L.;
 RT "Primary structure of the adult human skeletal muscle voltage-

RT dependent sodium channel.";
 RL Ann. Neurol. 31:131-137(1992).
 RN [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=92134303; PubMed=1310396;
 RA Wang J., Rojas C.V., Zhou J., Schwartz L.S., Nicholas H.,
 RA Hoffmann E.P.;
 RT "Sequence and genomic structure of the human adult skeletal muscle
 RT sodium channel alpha subunit gene on 17q.";
 RL Biochem. Biophys. Res. Commun. 182:794-801(1992).
 RN [3]
 RP SEQUENCE FROM N.A., VARIANT MYASTHENIC SYNDROME GLU-1442, AND VARIANTS
 RP LEU-246, ASP-559 AND ASN-1376.
 RX MEDLINE=22684480; PubMed=12766226; DOI=10.1073/pnas.1230273100;
 RA Teujino A., Maertens C., Ono K., Shen X.-M., Fukuda T., Harper C.M.,
 RA Cannon S.C., Engel A.G.;
 RT "Myasthenic syndrome caused by mutation of the SCN4A sodium channel";
 RT Proc. Natl. Acad. Sci. U.S.A. 100:7377-7382(2003).
 RN [4]
 RP SEQUENCE FROM N.A., AND VARIANT ASN-1376.
 RX MEDLINE=93338444; PubMed=1339144;
 RA McClatchey A.I., Lin C.S., Wang J., Hoffman E.P., Rojas C.V.,
 RA Gusella J.F.;
 RT "The genomic structure of the human skeletal muscle sodium channel
 RT gene.";
 RL Hum. Mol. Genet. 1:521-521(1992).
 RN [5]
 RP SEQUENCE OF 1305-1339 FROM N.A., AND VARIANTS PMC VAL-1306 AND
 RP MET-1313.
 RX MEDLINE=92514689; PubMed=1310898;
 RA McClatchey A.I., van den Berg P., Pericak-Vance M.A., Raekind W.,
 RA Verellen C., McKenna-Yasek D., Rao K., Haines J.L., Bird T.,
 RA Brown R.H. Jr., Gusella J.F.;
 RT "Temperature-sensitive mutations in the III-IV cytoplasmic loop region
 RT of the skeletal muscle sodium channel gene in paramyotonia
 RT congenita.";
 RL Cell 68:769-774(1992).
 RN [6]
 RP VARIANT HYP MET-704.
 RX MEDLINE=92069747; PubMed=1659948;
 RA Ptacek L.J., George A.L. Jr., Griggs R.C., Tawil R., Kallen R.G.,
 RA Barchi R.L., Robertson M., Leppert M.F.;
 RT "Identification of a mutation in the gene causing hyperkalemic
 RT periodic paralysis.";
 RL Cell 67:1021-1027(1991).
 RN [7]
 RP VARIANT HYP VAL-1592.
 RX MEDLINE=92065978; PubMed=1659668;
 RA Rojas C.V., Wang J., Schwartz L.S., Hoffman E.P., Powell B.R.,
 RA Brown R.H. Jr.;
 RT "A Met-to-Val mutation in the skeletal muscle Na+ channel alpha-
 RT subunit in hyperkalemic periodic paralysis.";
 RL Nature 354:387-389(1991).
 RN [8]
 RP VARIANTS PMC PHE-804 AND THR-1156.
 RX MEDLINE=93265141; PubMed=1338909;
 RA McClatchey A.I., McKenna-Yasek D., Cros D., Worthen H.G., Kuncel R.W.,
 RA Desilva S.M., Corbiach D.R., Gusella J.F., Brown R.H. Jr.;
 RT "Novel mutations in families with unusual and variable disorders of
 RT the skeletal muscle sodium channel.";
 RL Nat. Genet. 2:148-152(1992).
 RN [9]
 RP VARIANTS PMC CYS-1448 AND HIS-1448.
 RX MEDLINE=92265302; PubMed=1316765;
 RA Ptacek L.J., George A.L. Jr., Barchi R.L., Griggs R.C., Riggs J.E.,
 RA Robertson M., Leppert M.F.;
 RT "Mutations in an 84 segment of the adult skeletal muscle sodium
 RT channel cause paramyotonia congenita.";
 RL Neuron 8:891-897(1992).
 RN [10]
 RP VARIANT PMC ARG-1433.
 RX MEDLINE=93270429; PubMed=8388676;
 RA Ptacek L.J., Gouw L., Kwiatkowski H., McManis P., Mendell J.R.,

RA Barohn R.J., George A.L. Jr., Barchi R.L., Robertson M., Leppert M.F.;
 RT "Sodium channel mutations in paramyotonia congenita and hypokalemic
 RT periodic paralysis.";
 RL Ann. Neurol. 33:300-307(1993).
 RN [11]
 RP VARIANTS PMC ALA-1306; GLU-1306 AND VAL-1306.
 RX MEDLINE=94141728; PubMed=8308722;
 RA Lerche H., Heine R., Pika U., George A.L. Jr., Mitrovic N.,
 RA Browatzki M., Weiss T., Rivet-Bastide M., Franke C., Lomomaco M.,
 RA Ricker K., Lehmann-Horn F.;
 RT "Human sodium channel myotonia: slowed channel inactivation due to
 RT substitutions for a glycine within the III-IV linker.";
 RL J. Physiol. (Lond.) 470:13-22(1993).
 RN [12]
 RP VARIANT PMC MET-1589.
 RX MEDLINE=94061027; PubMed=8242056;
 RA Heine R., Pika U., Lehmann-Horn F.;
 RT "A novel SCN4A mutation causing myotonia aggravated by cold and
 RT potassium.";
 RL Hum. Mol. Genet. 2:1349-1353(1993).
 RN [13]
 RP VARIANT POTASSIUM-AGGRAVATED MYOTONIA VAL-1160.
 RX PubMed=8058156;
 RA Ptacek L.J., Tawil R., Griggs R.C., Meola G., McManis P., Barohn R.J.,
 RA Mendell J.R., Harris C., Spitzer R., Santiago F., Leppert M.F.;
 RT "Sodium channel mutations in acetazolamide-responsive myotonia
 RT congenita, paramyotonia congenita, and hyperkalemic periodic
 RT paralysis.";
 RL Neurology 44:1500-1503(1994).
 RN [14]
 RP VARIANT PARAMYOTONIA WITHOUT COLD PARALYSIS ILE-1293.
 RX MEDLINE=96154961; PubMed=8580427;
 RA Koch M.C., Baumbach K., George A.L., Ricker K.;
 RT "Paramyotonia congenita without paralysis on exposure to cold: a novel
 RT mutation in the SCN4A gene (Val1293Ile).";
 RL NeuroReport 6:2001-2004(1995).
 RN [15]
 RP VARIANT POTASSIUM-AGGRAVATED MYOTONIA MET-445.
 RX PubMed=9392583;
 RA Rosenfeld J., Sloan-Brown K., George A.L. Jr.;
 RT "A novel muscle sodium channel mutation causes painful congenital
 RT myotonia.";
 RL Ann. Neurol. 42:811-814(1997).
 RN [16]
 RP VARIANT POTASSIUM-AGGRAVATED MYOTONIA MET-445.
 RX PubMed=10218481;
 RA Wang D.W., Vanneccarr D., Ruben P.C., George A.L. Jr., Bennett P.B.;
 RT "Functional consequences of a domain I/S6 segment sodium channel
 RT mutation associated with painful congenital myotonia.";
 RL FEBS Lett. 448:231-234(1999).
 RN [17]
 RP VARIANT HYPOKPP HIS-669.
 RX PubMed=10599760;
 RA Bulman D.E., Scoggan K.A., van Oene M.D., Nicolle M.W., Hahn A.F.,
 RA Toller L.L., Ebers G.C.;
 RT "A novel sodium channel mutation in a family with hypokalemic periodic
 RT paralysis.";
 RL Neurology 53:1932-1936(1999).
 RN [18]
 RP VARIANT HYPOKPP SER-1158.
 RX PubMed=10851391;
 RA Sugita Y., Aoki T., Sugiyama Y., Hida C., Ogata M., Yamamoto T.;
 RT "Temperature-sensitive sodium channelopathy with heat-induced myotonia
 RT and cold-induced paralysis.";
 RL Neurology 54:2179-2181(2000).
 RN [19]
 RP VARIANTS HYPOKPP GLY-672 AND HIS-672.
 RX PubMed=10944223;
 RA Jurkat-Rott K., Mitrovic N., Hang C., Kouzmekine A., Ializzo P.,
 RA Herzog J., Lerche H., Nicole S., Vale-Santos J., Chauveau D.,
 RA Fontaine B., Lehmann-Horn F.;
 RT "Voltage-sensor sodium channel mutations cause hypokalemic periodic
 RT paralysis type 2 by enhanced inactivation and reduced current.";

RL Proc. Natl. Acad. Sci. U.S.A. 97:9549-9554(2000).
 RN [20]
 RP VARIANT HYPOKPP SER-672.
 RX PubMed=11558801;
 RA Bendahhou S., Cummins T.R., Griggs R.C., Fu Y.H., Ptacek L.J.;
 RT "Sodium channel inactivation defects are associated with
 RT acetazolamide-exacerbated hypokalemic periodic paralysis.";
 RL Ann. Neurol. 50:417-420(2001).
 RN [21]
 RP VARIANT HYPOKPP SER-672.
 RX PubMed=11591859;
 RA Davies N.P., Ruseon L.H., Samuel M., Hanna M.G.;
 RT "Sodium channel gene mutations in hypokalemic periodic paralysis: an
 RT uncommon cause in the UK.";
 RL Neurology 57:1323-1325(2001).
 CC -1- FUNCTION: This protein mediates the voltage-dependent sodium ion
 CC permeability of excitable membranes. Assuming opened or closed
 CC conformations in response to the voltage difference across the
 CC membrane, the protein forms a sodium-selective channel through
 CC which Na+ ions may pass in accordance with their electrochemical
 CC gradient. This sodium channel may be present in both denervated
 CC and innervated skeletal muscle.
 CC -1- SUBUNIT: Muscle sodium channels contain an alpha subunit and a
 CC smaller beta subunit. Interacts with the PDZ domain of the
 CC synaptobrevin SNCA1, SNCA1 and SNCA2 (by similarity).
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
 CC -1- DOMAIN: The sequence contains 4 internal repeats, each with 5
 CC hydrophobic segments (S1,S2,S3,S5,S6) and one positively charged
 CC segment (S4). Segments S4 are probably the voltage-sensors and are
 CC characterized by a series of positively charged amino acids at
 CC every third position.
 CC -1- DISEASE: Defects in SCN4A are the cause of paramyotonia congenita
 CC of von Eulenburg (PMC) [MIM:168300]. PMC is an autosomal dominant
 CC sodium channel disease characterized by myotonia, increased by
 CC exposure to cold, intermittent flaccid paresis, not necessarily

Query Match 71.4%; Score 20; DB 1; Length 1836;
 Best Local Similarity 100.0%; Pred. No. 2.6e-10;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 9 IFGSPFLNLLIIVAVMAY 28
 DB 431 IFGSPFLNLLIIVAVMAY 450

RESULT 18
 CIN4 RAT STANDARD; PRT; 1840 AA.
 AC P15390;
 DT 01-APR-1990 (Rel. 14, Created)
 DT 01-APR-1990 (Rel. 14, Last sequence update)
 DT 05-JUL-2004 (Rel. 44, Last annotation update)
 DE Sodium channel protein type IV alpha subunit (Voltage-gated sodium
 DE channel alpha subunit Nav1.4) (Sodium channel protein, skeletal muscle
 DE alpha-subunit) (Mn-1).
 GN Name=Scn4a;
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 OC NCBI_TaxID=10116;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=90148778; PubMed=2559760;
 RA Tytner J.S., Cooperman S.S., Tomiko S.A., Zhou J., Crean S.M.,
 RA Boyle M.B., Kallen R.G., Sheng Z., Barchi R.L., Sigworth F.J.,
 RA Goodman R.H., Agnew W.S., Mandel G.;
 RT "Primary structure and functional expression of a mammalian skeletal
 RT muscle sodium channel.";
 RL Neuron 3:33-49(1989).
 CC -1- FUNCTION: This protein mediates the voltage-dependent sodium ion
 CC permeability of excitable membranes. Assuming opened or closed
 CC conformations in response to the voltage difference across the
 CC membrane, the protein forms a sodium-selective channel through

CC which Na+ ions may pass in accordance with their electrochemical
 CC gradient. This sodium channel may be present in both denervated
 CC and innervated skeletal muscle.
 CC -1- SUBUNIT: Muscle sodium channels contain an alpha subunit and a
 CC smaller beta subunit. Interacts with the PDZ domain of the
 CC syntrophin SYNT1, SYNT1 and SYNT2 (By similarity).
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
 CC -1- DOMAIN: The sequence contains 4 internal repeats, each with 5
 CC hydrophobic segments (S1, S2, S3, S5, S6) and one positively charged
 CC segment (S4). Segments S4 are probably the voltage-sensors and are
 CC characterized by a series of positively charged amino acids at
 CC every third position.
 CC -1- SIMILARITY: Belongs to the sodium channel family.
 CC -1- SIMILARITY: Contains 1 IQ domain.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL, M26643; AAA41682.1; -.
 CC PIR, JN0007; CHRTM1.
 CC HSSP, P04775; IBYV.
 DR InterPro: IPR001582; Ca/Na_pore.
 DR InterPro: IPR002111; Cat_channel_TrpL.
 DR InterPro: IPR005821; Ion_trans.
 DR InterPro: IPR000048; IQ_region.
 DR InterPro: IPR005820; M-channel_nlg.
 DR InterPro: IPR001696; Na_channel1.
 DR InterPro: IPR008052; Na_channel4.
 DR InterPro: IPR010526; Na_trans_assoc.
 DR Pfam: PF00520; Ion_trans_4.
 DR Pfam: PF00612; IQ_1.
 DR Pfam: PF06512; Na_trans_assoc_1.
 DR PRINTS: PR00170; NACHANNEL.
 DR PRINTS: PR01665; NACHANNEL4.
 DR PROSITE: PS50096; IQ_1.
 KW Glycoprotein; Ion transport; Ionic channel; Multigene family;
 KW Phosphorylation; Repeat; Sodium channel; Transmembrane;
 KW Voltage-gated channel.
 FT TRANSMEM 130 152 S1 of repeat I.
 FT TRANSMEM 156 179 S2 of repeat I.
 FT TRANSMEM 192 212 S3 of repeat I.
 FT TRANSMEM 214 233 S4 of repeat I.
 FT TRANSMEM 252 274 S5 of repeat I.
 FT TRANSMEM 417 444 S6 of repeat I.
 FT TRANSMEM 571 593 S1 of repeat II.
 FT TRANSMEM 663 685 S2 of repeat II.
 FT TRANSMEM 696 721 S3 of repeat II.
 FT TRANSMEM 722 737 S4 of repeat II.
 FT TRANSMEM 756 778 S5 of repeat II.
 FT TRANSMEM 832 859 S6 of repeat II.
 FT TRANSMEM 1084 1105 S1 of repeat III.
 FT TRANSMEM 1117 1140 S2 of repeat III.
 FT TRANSMEM 1149 1168 S3 of repeat III.
 FT TRANSMEM 1195 1215 S4 of repeat III.
 FT TRANSMEM 1215 1236 S5 of repeat III.
 FT TRANSMEM 1324 1351 S6 of repeat III.
 FT TRANSMEM 1405 1427 S1 of repeat IV.
 FT TRANSMEM 1437 1460 S2 of repeat IV.
 FT TRANSMEM 1468 1487 S3 of repeat IV.
 FT TRANSMEM 1502 1522 S4 of repeat IV.
 FT TRANSMEM 1535 1556 S5 of repeat IV.
 FT TRANSMEM 1627 1653 S6 of repeat IV.
 FT DOMAIN 1720 1749 IQ.
 FT CARBOHYD 288 288 N-linked (GlcNAc...) (potential).
 FT CARBOHYD 291 291 N-linked (GlcNAc...) (potential).
 FT CARBOHYD 297 297 N-linked (GlcNAc...) (potential).
 FT CARBOHYD 303 303 N-linked (GlcNAc...) (potential).
 FT CARBOHYD 309 309 N-linked (GlcNAc...) (potential).

FT CARBOHYD 315 315 N-linked (GlcNAc...) (potential).
 FT CARBOHYD 327 327 N-linked (GlcNAc...) (potential).
 FT CARBOHYD 356 356 N-linked (GlcNAc...) (potential).
 FT CARBOHYD 502 502 N-linked (GlcNAc...) (potential).
 FT CARBOHYD 954 954 N-linked (GlcNAc...) (potential).
 FT CARBOHYD 1198 1198 N-linked (GlcNAc...) (potential).
 FT MOD RES 56 56 Phosphoserine (by PKA) (potential).
 FT MOD RES 251 251 Phosphoserine (by PKA) (potential).
 FT MOD RES 1321 1321 Phosphoserine (by PKA) (potential).
 FT MOD RES 1504 1504 Phosphoserine (by PKA) (potential).
 FT SEQUENCE 1840 AA; 208865 MW; CSDC09D93DD9FAD6 CRC64;
 SQ
 Query Match 71.4%; Score 20; DB 1; Length 1840;
 Best Local Similarity 100.0%; Pred. No. 2.6e-10;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 9 IFLGSFYILNLLAVVAMAY 28
 DB 425 IFLGSFYILNLLAVVAMAY 444
 RESULT 19
 070611 PRELIMINARY; PRT; 1840 AA.
 AC 070611;
 DT 01-AUG-1998 (TrEMBLrel. 07, Created)
 DT 01-AUG-1998 (TrEMBLrel. 07, Last sequence update)
 DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
 DE Rat skeletal muscle type I voltage-gated sodium channel (RSMK1)
 DE variant.
 GN Name=SCN4A;
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 OC Mammalia; Euteria; Rodentia; Sciurognathi; Muridae; Rattus.
 OX NCBI_TaxID=10116;
 RX [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=Copenhagen; TISSUE=Prostate;
 RX MEDLINE=98273645; PubMed=9613589;
 RA Disb J.K.J., Stewart D., Fraser S.P., Black J.A., Dibb-Hajj S.,
 RA Waxman S.G., Archer S.N., Djamgoz M.B.A.;
 RT "Expression of skeletal muscle-type voltage-gated Na+ channel in rat
 RT and human prostate cancer cell lines.";
 RL FEBS Lett. 427:5-10(1998).
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
 CC -1- SIMILARITY: Belongs to the sodium channel family.
 CC EMBL, Y17153; CAA76659.1; -.
 DR HSSP, P04775; IBYV.
 DR GO: GO:0016021; C:integral to membrane; IEA.
 DR GO: GO:0005158; C:voltage-gated sodium channel complex; IEA.
 DR GO: GO:0005246; F:cation channel activity; IEA.
 DR GO: GO:0005246; F:voltage-gated sodium channel activity; IEA.
 DR GO: GO:0006812; P:cation transport; IEA.
 DR GO: GO:0006814; P:sodium ion transport; IEA.
 DR InterPro: IPR001682; Ca/Na_pore.
 DR InterPro: IPR002111; Cat_channel_TrpL.
 DR InterPro: IPR005821; Ion_trans.
 DR InterPro: IPR000048; IQ_region.
 DR InterPro: IPR005820; M-channel_nlg.
 DR InterPro: IPR001696; Na_channel1.
 DR InterPro: IPR008052; Na_channel4.
 DR InterPro: IPR010526; Na_trans_assoc.
 DR Pfam: PF00520; Ion_trans_4.
 DR Pfam: PF00612; IQ_1.
 DR Pfam: PF06512; Na_trans_assoc_1.
 DR PRINTS: PR00170; NACHANNEL.
 DR PRINTS: PR01665; NACHANNEL4.
 DR SMART: SM00015; IQ_1.
 DR PROSITE: PS50096; IQ_1.
 DR Ion transport; Ionic channel; Sodium channel; Transmembrane;
 KW Transport; Voltage-gated channel.
 SQ SEQUENCE 1840 AA; 208823 MW; BIDPFA538E264B40 CRC64;

Query Match 71.4%; Score 20; DB 2; Length 1840;
 Best Local Similarity 100.0%; Pred. No. 2.6e-10;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 9 IFLGSFYLLINILIAVAVNAVY 28
 |||||
 Db 425 IFLGSFYLLINILIAVAVNAVY 444

RESULT 20
 QPER60
 ID QPER60 PRELIMINARY; PRT; 1841 AA.
 AC QPER60;
 DT 01-MAR-2001 (TREMBlrel. 16, Created)
 DT 01-MAR-2001 (TREMBlrel. 16, Last sequence update)
 DT 01-MAR-2004 (TREMBlrel. 26, Last annotation update)
 DE Voltage-gated sodium channel.
 GN Name=Scn4a;
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OK NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=balb/c; TISSUE=Heart;
 RX MEDLINE=21823196; PubMed=11834499;
 RA Zimmer T., Benndorf K.;
 RT "The mouse heart sodium channel (mH1): cloning and characterization of
 RT alternatively spliced variants."
 RL Am. J. Physiol. Heart Circ. Physiol. 282:H1007-H1017(2002).
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
 DR EMBL: A428787; CAC17146.1; -.
 DR HSSP: P04775; IBY.
 DR MGD; MGI:98250; Scn4a.
 DR GO; GO:0016021; C:integral to membrane; IEA.
 DR GO; GO:0001518; C:voltage-gated sodium channel complex; IEA.
 DR GO; GO:0005261; F:cation channel activity; IEA.
 DR GO; GO:0005248; F:voltage-gated sodium channel activity; IEA.
 DR GO; GO:0006814; P:sodium ion transport; IEA.
 DR GO; GO:0006814; P:sodium ion transport; IEA.
 DR InterPro: IPR001682; Ca/Na_pore.
 DR InterPro: IPR002111; Cat channel_Typl.
 DR InterPro: IPR005821; Ion trans.
 DR InterPro: IPR005820; M+channel_nlg.
 DR InterPro: IPR001686; Na_channel.
 DR InterPro: IPR008052; Na_channel4.
 DR InterPro: IPR010526; Na_trans_assoc.
 DR Pfam: PF00520; Ion_trans_4.
 DR Pfam: PF00612; IQ_1.
 DR Pfam: PF06512; Na_trans_assoc_1.
 DR PRINTS: PRO0170; NACHANNEL.
 DR PRINTS: PRO1665; NACHANNEL4.
 DR SMART: SM00015; IQ_1.
 DR PROSITE: PSS0096; IQ_1.
 KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
 KW Transport; Voltage-gated channel1.
 SQ SEQUENCE 1841 AA; 208196 MW; 0766DFD33A9E0B55 CRC64;

Query Match 71.4%; Score 20; DB 2; Length 1841;
 Best Local Similarity 100.0%; Pred. No. 2.6e-10;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 9 IFLGSFYLLINILIAVAVNAVY 28
 |||||
 Db 425 IFLGSFYLLINILIAVAVNAVY 444

RESULT 21
 Q090518
 AC Q090518; PRELIMINARY; PRT; 530 AA.
 RT

DT 01-NOV-1996 (TREMBlrel. 01, Created)
 DT 01-NOV-1996 (TREMBlrel. 01, Last sequence update)
 DT 01-JUN-2003 (TREMBlrel. 24, Last annotation update)
 DE Sodium channel alpha subunit (fragment).
 OS Fugu rubripes (Japanese pufferfish) (Takifugu rubripes).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
 OC Acanthomorpha; Acanthopterygii; Perciformes; Tetraodontiformes;
 OC Tetraodontidae; Tetraodontidae; Takifugu.
 OK NCBI_TaxID=31033;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Brain;
 RA Nakazawa A.;
 RL Submitted (Aug-1994) to the EMBL/GenBank/DBJ databases.
 DR EMBL: D37976; BAA07194.1; -.
 DR GO; GO:0016021; C:integral to membrane; IEA.
 DR GO; GO:0005261; F:cation channel activity; IEA.
 DR GO; GO:0006812; P:cation transport; IEA.
 DR InterPro: IPR005821; Ion trans.
 DR InterPro: IPR005820; M+channel_nlg.
 DR Pfam: PF00520; Ion_trans_1.
 KW Ion transport; Ionic channel; Transmembrane; Transport.
 FT NON TER 530 530
 SQ SEQUENCE 530 AA; 60164 MW; 2F9AB902C8F74071 CRC64;

Query Match 57.1%; Score 16; DB 2; Length 530;
 Best Local Similarity 100.0%; Pred. No. 6.2e-07;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 YMFVIVIFLGSFYLL 16
 |||||
 Db 392 YMFVIVIFLGSFYLL 407

RESULT 22
 Q080378
 ID Q080378 PRELIMINARY; PRT; 711 AA.
 AC Q080378;
 DT 01-JUN-2003 (TREMBlrel. 24, Created)
 DT 01-JUN-2003 (TREMBlrel. 24, Last sequence update)
 DT 01-OCT-2003 (TREMBlrel. 25, Last annotation update)
 DE Similar to sodium channel, voltage-gated, type 1, alpha
 DE polypeptide.
 GN Name=zgc:55600;
 OS Brachydanio rerio (Zebrafish) (Danio rerio).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;
 OC Cyprinidae; Danio.
 OK NCBI_TaxID=7955;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=AB; TISSUE=whole body;
 RX MEDLINE=22388257; PubMed=12477932;
 RA Strausberg R.L., Feingold B.A., Grouse L.H., Derge J.G.,
 RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
 RA Hopkins R.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
 RA Hootkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
 RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
 RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
 RA Brownstein M.O., Ushed T.B., Toshiyuki S., Carninci P., Prange C.,
 RA Raha S.S., Loquellano N.A., Peters G.J., Adamson R.D., Mullaly S.J.,
 RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
 RA Villalón D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
 RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Huik S.W.,
 RA Faley J., Helton E., Kettelman M., Madan A., Rodriguez S., Sanchez A.,
 RA Whiting M., Madan A., Young A.C., Suvchenko Y., Bouffard G.G.,
 RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M., Butterfield Y.S.,
 RA Krzywinski M.I., Skalski U., Smallus D.E., Schnerch A., Schein J.B.,
 RA Jones S.J., Mair M.A.;
 RT "Generation and initial analysis of more than 15,000 full-length human
 RT and mouse cDNA sequences.";


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RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=AB; TISSUE=whole body;
RA Strausberg R.;
DR Submitted (JAN-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC044197; AAH44197.1; -.
DR HSSP; P08104; I0G9.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0005261; P:cation channel activity; IEA.
DR GO; GO:0006812; P:cation channel activity; IEA.
DR InterPro; IPR0016821; Ca/Na_pore.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR005820; M+channel_nlg.
DR Pfam; PF00520; Ion_trans_1.
DR Ion transport; Ionic channel; Transmembrane; Transport.
KW SEQUENCE 711 AA; 80547 MW; 0BDE497E3BF85033 CRC64;

Query Match 57.1%; Score 16; DB 2; Length 711;
Best Local Similarity 100.0%; Pred. No. 7, 7e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 YMFVLYVIFLGSFYL 16
Db 382 YMFVLYVIFLGSFYL 397

RESULT 23
ID Q28371 PRELIMINARY; PRT; 1834 AA.
AC Q28371;
DT 01-NOV-1996 (TrEMBLrel. 01, Created)
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE Skeletal muscle sodium channel alpha-subunit.
OS Equus caballus (Horse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Perissodactyla; Equidae; Equus.
OX NCBI_Taxid=9796;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Skeletal muscle plasma membrane;
RA Stephan D.A., Wang J., Spier S., Hoffman E.P.;
RL Submitted (MAY-1995) to the EMBL/GenBank/DBJ databases.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
CC -1- SIMILARITY: Belongs to the sodium channel family.
DR EMBL; U25990; AAA67366.1; -.
DR HSSP; P04775; 1BYV.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0001518; C:voltage-gated sodium channel complex; IEA.
DR GO; GO:0005261; P:cation channel activity; IEA.
DR GO; GO:0005248; P:voltage-gated sodium channel activity; IEA.
DR GO; GO:0006812; P:cation transport; IEA.
DR GO; GO:0006814; P:sodium ion transport; IEA.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR002111; Cat_channel_TrpL.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IQ_region.
DR InterPro; IPR005820; M+channel_nlg.
DR InterPro; IPR001696; Na_channel.
DR InterPro; IPR008052; Na_channel.
DR InterPro; IPR010526; Na_trans_assoc.
DR InterPro; IPR001680; WD40.
DR Pfam; PF00520; Ion_trans_4.
DR Pfam; PF06512; IQ_1.
DR Pfam; PF06512; Na_trans_assoc; 1.
DR PRINTS; PR00170; NACHANNEL.
DR PRINTS; PR01665; NACHANNEL4.
DR SMART; SM00015; IQ_1.
DR PROSITE; PS00096; IQ_1.
DR PROSITE; PS00678; WD_REPEATS_1; UNKNOWN_1.
DR Ion transport; Ionic channel; Sodium channel; Transmembrane;
KW Transport; Voltage-gated channel.

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SQ SEQUENCE 1834 AA; 207485 MW; 01D62B25CD577D97 CRC64;
Query Match 57.1%; Score 16; DB 2; Length 1834;
Best Local Similarity 100.0%; Pred. No. 1, 6e-06;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 9 IFKGSFYLLNLLAVV 24
Db 425 IFKGSFYLLNLLAVV 440

RESULT 24
ID CIN3 RAT STANDARD; PRT; 1951 AA.
AC P08104;
DT 01-AUG-1988 (Rel. 08, Created)
DT 01-AUG-1988 (Rel. 08, Last sequence update)
DT 01-OCT-2004 (Rel. 45, Last annotation update)
DE Sodium channel protein type III alpha subunit (Voltage-gated sodium
DE channel alpha subunit Nav1.3) (Sodium channel protein, brain III alpha
DE subunit) (Voltage-gated sodium channel subtype III).
GN Name=Scn3a;
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_Taxid=10116;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Miscar;
RX MEDLINE=88137594; PubMed=2449363;
RA Kayano T., Noda M., Flockerzi V., Takahashi H., Numa S.;
RT "Primary structure of rat brain sodium channel III deduced from the
RT cDNA sequence.";
RL FEBS Lett. 228:187-194(1988).
CC -1- FUNCTION: Mediates the voltage-dependent sodium ion permeability
CC of excitable membranes. Assuming opened or closed conformations in
CC response to the voltage difference across the membrane, the
CC protein forms a sodium-selective channel through which Na(+) ions
CC may pass in accordance with their electrochemical gradient.
CC -1- SUBUNIT: The sodium channel consists of a large polypeptide and 2-
CC 3 smaller ones. This sequence represents a large polypeptide.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
CC -1- DOMAIN: The sequence contains 4 internal repeats, each with 5
CC hydrophobic segments (S1-S2, S3-S5, S6) and one positively charged
CC segment (S4). Segments S4 are probably the voltage-sensors and are
CC characterized by a series of positively charged amino acids at
CC every third position.
CC -1- SIMILARITY: Belongs to the sodium channel family.
CC -1- SIMILARITY: Contains 1 IQ domain.
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation-
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CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@sib-sib.ch).
DR EMBL; Y00766; CAA68735.1; -.
DR PIR; S00320; S00320.
DR PDB; 1OG9; NMR; A=156-176.
DR RGD; 3635; Scn3a.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR002111; Cat_channel_TrpL.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IQ_region.
DR InterPro; IPR005820; M+channel_nlg.
DR InterPro; IPR001696; Na_channel.
DR InterPro; IPR010526; Na_trans_assoc.
DR Pfam; PF00520; Ion_trans_4.
DR Pfam; PF06512; IQ_1.
DR PRINTS; PR00170; NACHANNEL.

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DR PROSITE; PS50096; IQ; FALSE NEG.
KW 3D-structure; Glycoprotein; Ion transport; Ionic channel;
KW Multigene family; Repeat; Sodium channel; Transmembrane;
KW Voltage-gated channel.
FT REPEAT 110 455
FT REPEAT 693 965
FT REPEAT 1139 1450
FT REPEAT 1459 1757
FT TRANSMEM 124 147
FT TRANSMEM 156 175
FT TRANSMEM 189 207
FT TRANSMEM 214 233
FT TRANSMEM 249 273
FT TRANSMEM 401 426
FT TRANSMEM 706 730
FT TRANSMEM 742 765
FT TRANSMEM 774 793
FT TRANSMEM 800 820
FT TRANSMEM 836 856
FT TRANSMEM 910 935
FT TRANSMEM 1153 1176
FT TRANSMEM 1190 1215
FT TRANSMEM 1222 1243
FT TRANSMEM 1248 1269
FT TRANSMEM 1289 1310
FT TRANSMEM 1393 1419
FT TRANSMEM 1473 1496
FT TRANSMEM 1508 1531
FT TRANSMEM 1538 1561
FT TRANSMEM 1572 1593
FT TRANSMEM 1609 1631
FT TRANSMEM 1698 1722
FT CARBOHYD 211 211
FT CARBOHYD 290 290
FT CARBOHYD 296 296
FT CARBOHYD 302 302
FT CARBOHYD 307 307
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FT CARBOHYD 624 624
FT CARBOHYD 835 835
FT CARBOHYD 1002 1002
FT CARBOHYD 1019 1019
FT CARBOHYD 1085 1085
FT CARBOHYD 1317 1317
FT CARBOHYD 1331 1331
SQ SEQUENCE 1951 AA; 221385 MM; 745B5B51524BD10B CRC64;

Query Match 57.1%; Score 16; DB 1; Length 1951;
Best Local Similarity 100.0%; Pred. No. 1.7e-06;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 YMIFVYVIFLGSFY 16
DB 400 YMIFVYVIFLGSFY 415

RESULT 25
Q9C007 PRELIMINARY; PRT; 1951 AA.
AC Q9C007;
DT 01-JUN-2001 (TrEMBLrel. 17, Created)
DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE Voltage-gated sodium channel alpha subunit splice variant SCN3A-
DE s.
GN Name=SCN3A;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
OX NCBI_TaxId=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Jeong S.-Y., Goto J., Kanazawa I.;

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RL Submitted (JAN-2000) to the EMBL/GenBank/DBJ databases.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (by similarity).
CC -1- SIMILARITY: Belongs to the sodium channel family.
DR EMBL; AF225986; AAK00218.1; -.
DR HSRP; P04775; 1BY; -.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0001518; C:voltage-gated sodium channel complex; IEA.
DR GO; GO:0005261; F:cation channel activity; IEA.
DR GO; GO:0005248; F:voltage-gated sodium channel activity; IEA.
DR GO; GO:0006812; P:cation transport; IEA.
DR GO; GO:0006814; P:sodium ion transport; IEA.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR002111; Cat_channel_Tyrl.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IQ_region.
DR InterPro; IPR005820; M+channel_nlg.
DR InterPro; IPR010526; Na_channel.
DR InterPro; IPR010526; Na_trans_assoc.
DR Pfam; PF00612; Ion_trans_4.
DR Pfam; PF06512; IQ_1.
DR Pfam; PF06512; Na_trans_assoc; 1.
DR PRINTS; PR00170; NACHANNEL.
DR SMART; SM00015; IQ; 1.
DR PROSITE; PS50096; IQ; 1.
KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
KW Transport; Voltage-gated channel.
SQ SEQUENCE 1951 AA; 221517 MM; 99AD4C032CE124AB CRC64;

Query Match 57.1%; Score 16; DB 2; Length 1951;
Best Local Similarity 100.0%; Pred. No. 1.7e-06;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 YMIFVYVIFLGSFY 16
DB 400 YMIFVYVIFLGSFY 415

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Search completed: January 27, 2005, 17:51:30
Job time : 96.5 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 27, 2005, 17:36:50 ; Search time 22.5 Seconds
(without alignments)
82.529 Million cell updates/sec

Title: US-10-608-584-1

Perfect score: 28

Sequence: 1 YMIFFVLIVIGSFYILNLIIVAMAY 28

Scoring table: OLIGO

Gapop 60.0 , Gapext 60.0

Searched: 478139 seqs, 66318000 residues

Word size : 0

Total number of hits satisfying chosen parameters: 478139

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: listing first 100 summaries

Database :

Issued_Patents_AA:*
1: /cgn2_6/pdata/1/1aa/5A_COMB.pep:*
2: /cgn2_6/pdata/1/1aa/5B_COMB.pep:*
3: /cgn2_6/pdata/1/1aa/6A_COMB.pep:*
4: /cgn2_6/pdata/1/1aa/6B_COMB.pep:*
5: /cgn2_6/pdata/1/1aa/ECTUS_COMB.pep:*
6: /cgn2_6/pdata/1/1aa/Backfill1.pep:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	28	100.0	2005	3	US-08-836-325-7
2	28	100.0	2005	4	US-09-457-571-7
3	21	75.0	1835	3	US-08-836-325-15
4	21	75.0	1835	4	US-09-457-571-15
5	21	75.0	1835	3	US-08-836-325-16
6	21	75.0	1835	4	US-09-457-571-16
7	21	75.0	1977	4	US-09-919-039-367
8	21	75.0	1977	4	US-08-836-325-10
9	21	75.0	1984	4	US-09-457-571-10
10	21	75.0	1989	3	US-08-836-325-11
11	21	75.0	1989	3	US-08-836-325-12
12	21	75.0	1989	3	US-09-457-571-11
13	21	75.0	1989	4	US-09-457-571-12
14	20	71.4	1836	4	US-10-162-012-24
15	20	71.4	1836	4	US-09-562-737-81
16	16	57.1	1924	4	US-08-843-417-10
17	16	57.1	1924	4	US-09-527-013-10
18	16	57.1	1956	4	US-08-843-417-2
19	13	46.4	1956	4	US-09-527-013-2
20	13	46.4	1956	4	US-08-669-656A-2
21	13	46.4	1957	4	US-08-669-656A-8
22	13	46.4	1957	4	US-08-669-656A-6
23	13	46.4	1957	4	US-08-669-656A-6
24	12	42.9	1233	4	US-09-354-147C-7
25	12	42.9	1233	4	US-09-354-147C-8
26	12	42.9	1243	4	US-09-354-147C-8
27	12	42.9	1791	4	US-09-354-147C-42

28	11	39.3	1976	3	US-09-024-020B-9	Sequence 9, Appl
29	11	39.3	1976	3	US-09-425-043-9	Sequence 9, Appl
30	11	39.3	1978	3	US-09-024-020B-3	Sequence 3, Appl
31	11	39.3	1978	3	US-09-425-043-3	Sequence 3, Appl
32	11	39.3	1988	3	US-09-024-020B-4	Sequence 4, Appl
33	11	39.3	1988	3	US-09-425-043-4	Sequence 4, Appl
34	11	39.3	2016	4	US-09-634-920-4	Sequence 4, Appl
35	11	39.3	2016	4	US-09-514-907A-2	Sequence 2, Appl
36	11	39.3	2016	4	US-09-896-994-2	Sequence 2, Appl
37	11	39.3	2016	4	US-09-840-125-4	Sequence 4, Appl
38	11	32.1	1024	4	US-09-562-737-82	Sequence 82, Appl
39	9	32.1	1024	4	US-09-562-737-84	Sequence 84, Appl
40	9	32.1	1024	4	US-09-562-737-85	Sequence 85, Appl
41	9	32.1	1024	4	US-09-562-737-86	Sequence 86, Appl
42	9	32.1	1024	4	US-09-562-737-87	Sequence 87, Appl
43	9	32.1	1024	4	US-09-562-737-88	Sequence 88, Appl
44	9	32.1	1024	4	US-09-562-737-89	Sequence 89, Appl
45	9	32.1	1024	4	US-09-562-737-90	Sequence 90, Appl
46	9	32.1	1765	4	US-09-354-147C-2	Sequence 2, Appl
47	9	32.1	1765	4	US-09-354-147C-3	Sequence 3, Appl
48	9	32.1	1765	4	US-09-354-147C-5	Sequence 5, Appl
49	8	28.6	1820	3	US-07-998-289B-8	Sequence 8, Appl
50	8	28.6	2100	2	US-08-808-793-23	Sequence 23, Appl
51	8	28.6	2100	2	US-08-772-512A-19	Sequence 19, Appl
52	8	28.6	2104	2	US-08-808-793-4	Sequence 4, Appl
53	8	28.6	2104	2	US-08-772-512A-4	Sequence 4, Appl
54	8	28.6	2105	2	US-08-808-793-3	Sequence 3, Appl
55	8	28.6	2105	3	US-08-772-512A-3	Sequence 3, Appl
56	7	25.0	562	4	US-09-328-352-4694	Sequence 4694, Ap
57	6	21.4	42	1	US-08-118-270-311	Sequence 311, Ap
58	6	21.4	42	5	PCR-US93-08528-311	Sequence 311, Ap
59	6	21.4	43	1	US-08-118-270-312	Sequence 310, Ap
60	6	21.4	43	5	PCR-US93-08528-310	Sequence 310, Ap
61	6	21.4	44	1	US-08-118-270-312	Sequence 312, Ap
62	6	21.4	44	5	PCR-US93-08528-312	Sequence 312, Ap
63	6	21.4	46	1	US-08-118-270-313	Sequence 313, Ap
64	6	21.4	46	5	PCR-US93-08528-313	Sequence 313, Ap
65	6	21.4	48	1	US-08-118-270-314	Sequence 314, Ap
66	6	21.4	48	5	PCR-US93-08528-314	Sequence 314, Ap
67	6	21.4	82	4	US-09-134-000C-5571	Sequence 5571, Ap
68	6	21.4	86	1	US-07-847-743B-19	Sequence 19, Appl
69	6	21.4	86	1	US-08-456-201-19	Sequence 19, Appl
70	6	21.4	86	2	US-08-330-161-17	Sequence 17, Appl
71	6	21.4	86	2	US-08-456-241-19	Sequence 19, Appl
72	6	21.4	86	2	US-08-440-401-17	Sequence 17, Appl
73	6	21.4	86	2	US-08-419-878B-17	Sequence 17, Appl
74	6	21.4	86	3	US-09-173-480-17	Sequence 17, Appl
75	6	21.4	86	5	PCR-US92-04255A-19	Sequence 19, Appl
76	6	21.4	110	4	US-09-328-352-5892	Sequence 5892, Ap
77	6	21.4	129	4	US-09-270-767-43684	Sequence 43684, A
78	6	21.4	133	4	US-09-270-767-38939	Sequence 38939, A
79	6	21.4	133	4	US-09-270-767-54156	Sequence 54156, A
80	6	21.4	136	4	US-09-248-796A-22644	Sequence 22644, A
81	6	21.4	147	4	US-09-270-767-38096	Sequence 38096, A
82	6	21.4	147	4	US-09-270-767-53313	Sequence 53313, A
83	6	21.4	150	4	US-09-328-352-7806	Sequence 7806, Ap
84	6	21.4	159	3	US-08-796-792-2	Sequence 2, Appl
85	6	21.4	159	4	US-09-491-795-2	Sequence 2, Appl
86	6	21.4	184	4	US-09-134-000C-5724	Sequence 6724, Ap
87	6	21.4	194	4	US-09-489-039A-8286	Sequence 8286, Ap
88	6	21.4	200	3	US-09-134-001C-1099	Sequence 3099, Ap
89	6	21.4	208	1	US-07-935-309-2	Sequence 2, Appl
90	6	21.4	208	2	US-08-039-364-2	Sequence 2, Appl
91	6	21.4	208	3	US-08-718-904-5	Sequence 5, Appl
92	6	21.4	208	3	US-08-718-904-7	Sequence 7, Appl
93	6	21.4	208	3	US-09-181-974-2	Sequence 2, Appl
94	6	21.4	208	3	US-09-158-710-2	Sequence 2, Appl
95	6	21.4	208	3	US-09-518-950-2	Sequence 2, Appl
96	6	21.4	208	4	US-09-449-249-5	Sequence 5, Appl
97	6	21.4	208	4	US-09-449-249-7	Sequence 7, Appl
98	6	21.4	208	4	US-10-138-158-18	Sequence 18, Appl
99	6	21.4	240	4	US-09-489-039A-8113	Sequence 8113, Ap
100	6	21.4	261	4	US-09-522-714-6	Sequence 6, Appl

ALIGNMENTS

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RESULT 1
US-08-836-325-7
; Sequence 7, Application US/08836325
; Patent No. 6110672
; GENERAL INFORMATION:
; APPLICANT: Mandel, Gail
; APPLICANT: Halegoua, Simon
; APPLICANT: Borden, Laurence A.
; TITLE OF INVENTION: Peripheral Nervous System Specific
; TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
; TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
; TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
; NUMBER OF SEQUENCES: 19
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C
; STREET: 1100 New York Ave., N. W., Suite 600
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20005-3934
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; FILING DATE: 02-NOV-1995
; APPLICATION NUMBER: 08/836,325
; FILING DATE: 2-MAY-1997
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/14251
; FILING DATE: 02-NOV-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/482,401
; FILING DATE: 07-JUN-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/334,029
; FILING DATE: 02-NOV-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Ludwig, Steven R.
; REGISTRATION NUMBER: 36,203
; REFERENCE/DOCKET NUMBER: 0917.0240002
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-371-2600
; TELEFAX: 202-371-2540
; INFORMATION FOR SEQ ID NO: 7:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2005 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
US-08-836-325-7

Query Match      100.0%; Score 28; DB 3; Length 2005;
Best Local Similarity 100.0%; Pred. No. 2.3e-19;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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OY 1 YWIFVLVIFLGSFYLINILAVVMAY 28
DB 401 YWIFVLVIFLGSFYLINILAVVMAY 428

RESULT 2
US-09-457-571-7
; Sequence 7, Application US/09457571
; Patent No. 6703486

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; GENERAL INFORMATION:
; APPLICANT: Mandel, Gail
; APPLICANT: Halegoua, Simon
; APPLICANT: Borden, Laurence A.
; TITLE OF INVENTION: Peripheral Nervous System Specific
; TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
; TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
; TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
; NUMBER OF SEQUENCES: 19
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C
; STREET: 1100 New York Ave., N. W., Suite 600
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20005-3934
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; FILING DATE: 09-DEC-1999
; APPLICATION NUMBER: US/09/457,571
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/836,325
; FILING DATE: 02-MAY-1997
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/14251
; FILING DATE: 02-NOV-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/482,401
; FILING DATE: 07-JUN-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/334,029
; FILING DATE: 02-NOV-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Ludwig, Steven R.
; REGISTRATION NUMBER: 36,203
; REFERENCE/DOCKET NUMBER: 0917.0240003
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-371-2600
; TELEFAX: 202-371-2540
; INFORMATION FOR SEQ ID NO: 7:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2005 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
US-09-457-571-7

Query Match      100.0%; Score 28; DB 4; Length 2005;
Best Local Similarity 100.0%; Pred. No. 2.3e-19;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 YWIFVLVIFLGSFYLINILAVVMAY 28
DB 401 YWIFVLVIFLGSFYLINILAVVMAY 428

RESULT 3
US-08-836-325-15
; Sequence 15, Application US/08836325
; Patent No. 6110672
; GENERAL INFORMATION:
; APPLICANT: Mandel, Gail
; APPLICANT: Halegoua, Simon
; APPLICANT: Borden, Laurence A.
; TITLE OF INVENTION: Peripheral Nervous System Specific
; TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
; TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational

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TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
TITLE OF INVENTION: Theeocf
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C.
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/836,325
FILING DATE: 2-MAY-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917.0240002
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2540
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 15:
SEQUENCE CHARACTERISTICS:
LENGTH: 1835 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: not relevant
MOLECULE TYPE: protein
US-08-836-325-15

Query Match 75.0%; Score 21; DB 3; Length 1835;
Best Local Similarity 100.0%; Pred. No. 1.6e-12;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 8 VIFLSFYINILAVVMAY 28
Db 363 VIFLSFYINILAVVMAY 383

RESULT 4
US-09-457-571-15
Sequence 15, Application US/09457571
Patent No. 6703486
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Halegoua, Simon
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C.
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/457,571
FILING DATE: 09-DEC-1999
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/836,325
FILING DATE: 02-MAY-1997
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917.0240003
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2540
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 15:
SEQUENCE CHARACTERISTICS:
LENGTH: 1835 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: not relevant
MOLECULE TYPE: protein
US-09-457-571-15

Query Match 75.0%; Score 21; DB 4; Length 1835;
Best Local Similarity 100.0%; Pred. No. 1.6e-12;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 8 VIFLSFYINILAVVMAY 28
Db 363 VIFLSFYINILAVVMAY 383

RESULT 5
US-08-836-325-16
Sequence 16, Application US/08836325
Patent No. 6110672
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Halegoua, Simon
APPLICANT: Borden, Laurence A.
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C.
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/836,325
FILING DATE: 2-MAY-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917.0240002
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 16:
SEQUENCE CHARACTERISTICS:
LENGTH: 1969 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-836-325-16

Query Match
Best Local Similarity 75.0%; Score 21; DB 3; Length 1969;
Matches 21; Conservative 0; Pred. No. 1,7e-12;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 VFILGSFYILNLIILAVVAMAY 28
DB 385 VFILGSFYILNLIILAVVAMAY 405

RESULT 6
US-09-457-571-16
Sequence 16, Application US/09457571
Patent No. 6703486
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Halsegoud, Simon
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Thereof, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESSES:
ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/457,571
FILING DATE: 09-DEC-1999
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/836,325
FILING DATE: 02-MAY-1997
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:

APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917.0240003
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2540
TELEFAX: 202-371-2600
INFORMATION FOR SEQ ID NO: 16:
SEQUENCE CHARACTERISTICS:
LENGTH: 1969 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: linear
MOLECULE TYPE: protein
US-09-457-571-16

Query Match
Best Local Similarity 75.0%; Score 21; DB 4; Length 1969;
Matches 21; Conservative 0; Pred. No. 1,7e-12;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 VFILGSFYILNLIILAVVAMAY 28
DB 385 VFILGSFYILNLIILAVVAMAY 405

RESULT 7
US-09-976-594-757
Sequence 757, Application US/09976594
Patent No. 6673549
GENERAL INFORMATION:
APPLICANT: Furness, Michael
APPLICANT: Buchbinder, Jenny
TITLE OF INVENTION: GENES EXPRESSED IN C3A LIVER CELL CULTURES TREATED WITH STEROIDS
FILE REFERENCE: PA-0041 US
CURRENT APPLICATION NUMBER: US/09/976,594
PRIOR FILING DATE: 2001-10-12
PRIOR APPLICATION NUMBER: 60/240,409
NUMBER OF SEQ ID NOS: 1143
SOFTWARE: PERL Program
SEQ ID NO 757
LENGTH: 1977
TYPE: PRT
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: misc feature
OTHER INFORMATION: Incyte ID No. 6673549 1719478CD1
US-09-976-594-757

Query Match
Best Local Similarity 75.0%; Score 21; DB 4; Length 1977;
Matches 21; Conservative 0; Pred. No. 1,7e-12;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 VFILGSFYILNLIILAVVAMAY 28
DB 385 VFILGSFYILNLIILAVVAMAY 405

RESULT 8
US-09-919-039-367
Sequence 367, Application US/09919039
Patent No. 6727066
GENERAL INFORMATION:
APPLICANT: Kaser, Matthew R.
TITLE OF INVENTION: GENES EXPRESSED IN TREATED HUMAN C3A LIVER CELL CULTURES
FILE REFERENCE: PA-0035 US
CURRENT APPLICATION NUMBER: US/09/919,039
CURRENT FILING DATE: 2002-09-09

PRIOR APPLICATION NUMBER: 60/222,113
PRIOR FILING DATE: 2000-07-28
NUMBER OF SEQ ID NOS: 401
SOFTWARE: PERL Program
SEQ ID NO 367
LENGTH: 1977
TYPE: PRT
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: misc_feature
OTHER INFORMATION: incyte ID No. 6727066 1719478CD1
US-09-919-039-367

Query Match 75.0%; Score 21; DB 4; Length 1977;
Best Local Similarity 100.0%; Pred. No. 1.7e-12;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 VIFLGSFYILNLIILAVVAMAY 28
DB 385 VIFLGSFYILNLIILAVVAMAY 405

RESULT 9
US-08-836-325-10
Sequence 10, Application US/08836325
Patent No. 6110672
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Halegoua, Simon
TITLE OF INVENTION: Borden, Laurence A.
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
TITLE OF INVENTION: Thereof
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESSES:
ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/836,325
FILING DATE: 2-MAY-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917.0240002
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2540
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 10:
SEQUENCE CHARACTERISTICS:
LENGTH: 1984 amino acids
TYPE: amino acid
TOPOLOGY: linear

MOLECULE TYPE: protein
US-08-836-325-10

Query Match 75.0%; Score 21; DB 3; Length 1984;
Best Local Similarity 100.0%; Pred. No. 1.7e-12;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 VIFLGSFYILNLIILAVVAMAY 28
DB 385 VIFLGSFYILNLIILAVVAMAY 405

RESULT 10
US-09-457-571-10
Sequence 10, Application US/09457571
Patent No. 6703486
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Halegoua, Simon
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
TITLE OF INVENTION: Thereof
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESSES:
ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/457,571
FILING DATE: 09-DEC-1999
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/836,325
FILING DATE: 02-MAY-1997
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917.0240003
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 10:
SEQUENCE CHARACTERISTICS:
LENGTH: 1984 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-09-457-571-10

Query Match 75.0%; Score 21; DB 4; Length 1984;
Best Local Similarity 100.0%; Pred. No. 1.7e-12;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 VIFLGSFYILNLIILAVVAMAY 28

Db 385 VIFLGSFYILNIIILAVVAMAY 405

```
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RESULT 11
US-08-836-325-11
; Sequence 11, Application US/08836325
; Patent No. 6110672
; GENERAL INFORMATION:
; APPLICANT: Mandel, Gail
; APPLICANT: Halegoua, Simon
; APPLICANT: Borden, Laurence A.
; TITLE OF INVENTION: Peripheral Nervous System Specific
; TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
; TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
; TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
; NUMBER OF SEQUENCES: 19
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
; STREET: 1100 New York Ave., N. W., Suite 600
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20005-3934
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: IBM PC compatible
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/836,325
; FILING DATE: 2-MAY-1997
; CLASSIFICATION: 514
; PRIORITY APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/14251
; FILING DATE: 02-NOV-1995
; APPLICATION NUMBER: 08/482,401
; FILING DATE: 07-JUN-1995
; PRIORITY APPLICATION DATA:
; APPLICATION NUMBER: 08/334,029
; FILING DATE: 02-NOV-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Ludwig, Steven R.
; REGISTRATION NUMBER: 36,203
; REFERENCE/DOCKET NUMBER: 0917.0240002
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-371-2600
; TELEFAX: 202-371-2540
; INFORMATION FOR SEQ ID NO: 11:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1989 amino acids
; TYPE: amino acid
; STRANDEDNESS: not relevant
; TOPOLOGY: not relevant
; MOLECULE TYPE: protein
; US-08-836-325-11

Query Match 75.0%; Score 21; DB 3; Length 1989;
Best Local Similarity 100.0%; Pred. No. 1.7e-12;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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QY 8 VIFLGSFYILNIIILAVVAMAY 28

Db 385 VIFLGSFYILNIIILAVVAMAY 405

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|||||
RESULT 12
US-08-836-325-12
; Sequence 12, Application US/08836325
; Patent No. 6110672
; GENERAL INFORMATION:
```

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|||||
; APPLICANT: Mandel, Gail
; APPLICANT: Halegoua, Simon
; APPLICANT: Borden, Laurence A.
; TITLE OF INVENTION: Peripheral Nervous System Specific
; TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
; TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
; TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
; NUMBER OF SEQUENCES: 19
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
; STREET: 1100 New York Ave., N. W., Suite 600
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20005-3934
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: IBM PC compatible
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/836,325
; FILING DATE: 2-MAY-1997
; CLASSIFICATION: 514
; PRIORITY APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/14251
; FILING DATE: 02-NOV-1995
; APPLICATION NUMBER: 08/482,401
; FILING DATE: 07-JUN-1995
; PRIORITY APPLICATION DATA:
; APPLICATION NUMBER: 08/334,029
; FILING DATE: 02-NOV-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Ludwig, Steven R.
; REGISTRATION NUMBER: 36,203
; REFERENCE/DOCKET NUMBER: 0917.0240002
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-371-2600
; TELEFAX: 202-371-2540
; INFORMATION FOR SEQ ID NO: 12:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1989 amino acids
; TYPE: amino acid
; STRANDEDNESS: not relevant
; TOPOLOGY: not relevant
; MOLECULE TYPE: protein
; US-08-836-325-12
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QY 8 VIFLGSFYILNIIILAVVAMAY 28

Db 385 VIFLGSFYILNIIILAVVAMAY 405

Query Match 75.0%; Score 21; DB 3; Length 1989;
Best Local Similarity 100.0%; Pred. No. 1.7e-12;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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RESULT 13
US-09-457-571-11
; Sequence 11, Application US/09457571
; Patent No. 6703486
; GENERAL INFORMATION:
; APPLICANT: Mandel, Gail
; APPLICANT: Halegoua, Simon
; APPLICANT: Borden, Laurence A.
; TITLE OF INVENTION: Peripheral Nervous System Specific
; TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
; TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
; TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
; NUMBER OF SEQUENCES: 19
; CORRESPONDENCE ADDRESS:
```


ADDRESSEE: STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C.
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/457,571
FILING DATE: 09-DEC-1999
CLASSIFICATION: 514
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: 08/836,325
FILING DATE: 02-MAY-1997
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917.0240003
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 11:
SEQUENCE CHARACTERISTICS:
LENGTH: 1989 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: not relevant
MOLECULE TYPE: protein
US-09-457-571-11

Query Match 75.0%; Score 21; DB 4; Length 1989;
Best Local Similarity 100.0%; Pred.No.1.7e-12;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 VIFLGSFYILNLIILAVNAY 28
DB 385 VIFLGSFYILNLIILAVNAY 405

RESULT 14
US-09-457-571-12
Sequence 12, Application US/09457571
Patent No. 6703486
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
TITLE OF INVENTION: Thereof
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C.
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/457,571
FILING DATE: 09-DEC-1999
CLASSIFICATION: 514
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: 08/836,325
FILING DATE: 02-MAY-1997
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917.0240003
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 12:
SEQUENCE CHARACTERISTICS:
LENGTH: 1989 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: not relevant
MOLECULE TYPE: protein
US-09-457-571-12

Query Match 75.0%; Score 21; DB 4; Length 1989;
Best Local Similarity 100.0%; Pred.No.1.7e-12;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 VIFLGSFYILNLIILAVNAY 28
DB 385 VIFLGSFYILNLIILAVNAY 405

RESULT 15
US-10-162-012-24
Sequence 24, Application US/10162012
Patent No. 6682597
GENERAL INFORMATION:
APPLICANT: Curtiss, Roy A.J.
APPLICANT: Siles-Santiago, Immaculada
TITLE OF INVENTION: NOVEL HUMAN ION CHANNEL AND TRANSPORTER FAMILY MEMBERS
FILE REFERENCE: 10448-190001
CURRENT APPLICATION NUMBER: US/10/162,012
CURRENT FILING DATE: 2002-06-04
PRIORITY APPLICATION NUMBER: US 60/209,845
PRIORITY FILING DATE: 2000-06-06
PRIORITY APPLICATION NUMBER: US 09/875,321
PRIORITY FILING DATE: 2001-06-06
PRIORITY APPLICATION NUMBER: PCT/US01/18340
PRIORITY FILING DATE: 2001-06-06
PRIORITY APPLICATION NUMBER: US 60/209,257
PRIORITY FILING DATE: 2000-06-05
PRIORITY APPLICATION NUMBER: US 09/875,423
PRIORITY FILING DATE: 2001-06-05
PRIORITY APPLICATION NUMBER: PCT/US01/18398
PRIORITY FILING DATE: 2001-06-05
PRIORITY APPLICATION NUMBER: US 60/209,238
PRIORITY FILING DATE: 2000-06-05
PRIORITY APPLICATION NUMBER: US 09/875,363
PRIORITY FILING DATE: 2001-06-05

RESULT 17

RESULT 18
 US-09-537-013-10
 Sequence 10, Application US/09527013
 Patent No. 6479259
 GENERAL INFORMATION:
 APPLICANT: Herman, Ronald C
 Delgado, Stephen G
 Fish, Linda M
 Sangameswaran, Lakshmi
 Rabert, Douglas K
 TITLE OF INVENTION: CLONED PERIPHERAL NERVE
 TETRODOTOXIN-RESISTANT SODIUM CHANNEL alpha-SUBUNIT
 NUMBER OF SEQUENCES: 10
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Heller, Herman White & McLaughlin
 STREET: 525 University Ave
 CITY: Palo Alto
 STATE: CA

COUNTRY: U.S.A.
ZIP: 94301
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/527,013
FILING DATE: 16-Mar-2000
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/843,417
FILING DATE: <Unknown>
ATTORNEY/AGENT INFORMATION:
NAME: Schmonsees, William
REGISTRATION NUMBER: 31,796
REFERENCE/DOCKET NUMBER: 28340-P1
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415)-324-7041
TELEFAX: (415)-324-0638
INFORMATION FOR SEQ ID NO: 10:
SEQUENCE CHARACTERISTICS:
LENGTH: 1956 amino acids
TYPE: amino acid
STRANDEDNESS: unknown
TOPOLOGY: unknown
MOLECULE TYPE: peptide
HYPOTHETICAL: NO
ANTI-SENSE: NO
SEQUENCE DESCRIPTION: SEQ ID NO: 10:
US-09-527-013-10

Query Match 57.1%; Score 16; DB 4; Length 1956;
Best Local Similarity 100.0%; Pred. No. 1.4e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 YMFPLVIFLGSFYL 16
DB 373 YMFPLVIFLGSFYL 388

RESULT 19
US-08-843-417-2
Sequence 2, Application US/08843417
Patent No. 6184349
GENERAL INFORMATION:
APPLICANT: Herman, Ronald C
APPLICANT: Delgado, Stephen G
APPLICANT: Fish, Linda M
APPLICANT: Sangameswaran, Lakshmi
APPLICANT: Rabert, Douglas K
TITLE OF INVENTION: CLONED PERIPHERAL NERVE
TITLE OF INVENTION: TETRODOTOXIN-RESISTANT SODIUM CHANNEL alpha-SUBUNIT
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: Heller Ehrman White & McCauliffe
STREET: 525 University Ave
CITY: Palo Alto
STATE: CA
COUNTRY: U.S.A.
ZIP: 94301
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/843,417
FILING DATE: April 15, 1997
CLASSIFICATION: 536
ATTORNEY/AGENT INFORMATION:
NAME: Schmonsees, William

REGISTRATION NUMBER: 31,796
REFERENCE/DOCKET NUMBER: 28340-P1
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415)-324-7041
TELEFAX: (415)-324-0638
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 1956 amino acids
TYPE: amino acid
STRANDEDNESS:
TOPOLOGY: not relevant
MOLECULE TYPE: protein
HYPOTHETICAL: YES
ORIGINAL SOURCE:
ORGANISM: rat
TISSUE TYPE: dorsal root ganglia
CELL TYPE: peripheral nerve
US-08-843-417-2

Query Match 46.4%; Score 13; DB 3; Length 1956;
Best Local Similarity 100.0%; Pred. No. 0.00012;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 FFVLVIFLGSFYL 16
DB 375 FFVLVIFLGSFYL 387

RESULT 20
US-09-527-013-2
Sequence 2, Application US/09527013
Patent No. 6479259
GENERAL INFORMATION:
APPLICANT: Herman, Ronald C
APPLICANT: Delgado, Stephen G
APPLICANT: Fish, Linda M
APPLICANT: Sangameswaran, Lakshmi
APPLICANT: Rabert, Douglas K
TITLE OF INVENTION: CLONED PERIPHERAL NERVE
TITLE OF INVENTION: TETRODOTOXIN-RESISTANT SODIUM CHANNEL alpha-SUBUNIT
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: Heller Ehrman White & McCauliffe
STREET: 525 University Ave
CITY: Palo Alto
STATE: CA
COUNTRY: U.S.A.
ZIP: 94301
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/527,013
FILING DATE: 16-Mar-2000
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/843,417
FILING DATE: <Unknown>
ATTORNEY/AGENT INFORMATION:
NAME: Schmonsees, William
REGISTRATION NUMBER: 31,796
REFERENCE/DOCKET NUMBER: 28340-P1
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415)-324-7041
TELEFAX: (415)-324-0638
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 1956 amino acids
TYPE: amino acid
STRANDEDNESS: <Unknown>
TOPOLOGY: not relevant

MOLECULE TYPE: protein
 HYPOTHETICAL: YES
 ORIGINAL SOURCE:
 ORGANISM: rat
 TISSUE TYPE: dorsal root ganglia
 CELL TYPE: peripheral nerve
 SEQUENCE DESCRIPTION: SEQ ID NO: 2:
 US-09-527-013-2

Query Match 46.4%; Score 13; DB 4; Length 1956;
 Best Local Similarity 100.0%; Pred. No. 0.00012;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 FFVLVIFLGSFY 16
 DB 375 FFVLVIFLGSFY 387

RESULT 21
 US-08-669-656A-2
 Sequence 2, Application US/08669656A
 Patent No. 6451554
 GENERAL INFORMATION:
 APPLICANT: Wood, John N.
 APPLICANT: Akopian, Armen N.
 TITLE OF INVENTION: Ion Channel
 NUMBER OF SEQUENCES: 31
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: ZENECA Pharmaceuticals
 STREET: 1800 Concord Pike, P.O. Box 15437
 CITY: Wilmington
 STATE: Delaware
 COUNTRY: USA
 ZIP: 19850
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: Patentin Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/669,656A
 FILING DATE: 24-JUN-1996
 CLASSIFICATION: 536
 ATTORNEY/AGENT INFORMATION:
 NAME: Hohenschutz, Liza D.
 REGISTRATION NUMBER: 33,712
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (302) 886-7466
 INFORMATION FOR SEQ ID NO: 2:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 1957 amino acids
 TYPE: amino acid
 TOPOLOGY: linear
 MOLECULE TYPE: protein
 US-08-669-656A-2

Query Match 46.4%; Score 13; DB 4; Length 1957;
 Best Local Similarity 100.0%; Pred. No. 0.00012;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 FFVLVIFLGSFY 16
 DB 375 FFVLVIFLGSFY 387

RESULT 22
 US-08-669-656A-8
 Sequence 8, Application US/08669656A
 Patent No. 6451554
 GENERAL INFORMATION:
 APPLICANT: Wood, John N.
 APPLICANT: Akopian, Armen N.

TITLE OF INVENTION: Ion Channel
 NUMBER OF SEQUENCES: 31
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: ZENECA Pharmaceuticals
 STREET: 1800 Concord Pike, P.O. Box 15437
 CITY: Wilmington
 STATE: Delaware
 COUNTRY: USA
 ZIP: 19850
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: Patentin Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/669,656A
 FILING DATE: 24-JUN-1996
 CLASSIFICATION: 536
 ATTORNEY/AGENT INFORMATION:
 NAME: Hohenschutz, Liza D.
 REGISTRATION NUMBER: 33,712
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (302) 886-7466
 INFORMATION FOR SEQ ID NO: 8:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 1957 amino acids
 TYPE: amino acid
 TOPOLOGY: linear
 MOLECULE TYPE: protein
 US-08-669-656A-8

Query Match 46.4%; Score 13; DB 4; Length 1957;
 Best Local Similarity 100.0%; Pred. No. 0.00012;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 FFVLVIFLGSFY 16
 DB 375 FFVLVIFLGSFY 387

RESULT 23
 US-08-669-656A-6
 Sequence 6, Application US/08669656A
 Patent No. 6451554
 GENERAL INFORMATION:
 APPLICANT: Wood, John N.
 APPLICANT: Akopian, Armen N.
 TITLE OF INVENTION: Ion Channel
 NUMBER OF SEQUENCES: 31
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: ZENECA Pharmaceuticals
 STREET: 1800 Concord Pike, P.O. Box 15437
 CITY: Wilmington
 STATE: Delaware
 COUNTRY: USA
 ZIP: 19850
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: Patentin Release #1.0, Version #1.25
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/669,656A
 FILING DATE: 24-JUN-1996
 CLASSIFICATION: 536
 ATTORNEY/AGENT INFORMATION:
 NAME: Hohenschutz, Liza D.
 REGISTRATION NUMBER: 33,712
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (302) 886-7466
 INFORMATION FOR SEQ ID NO: 6:

SEQUENCE CHARACTERISTICS:
 LENGTH: 2132 amino acids
 TYPE: amino acid
 TOPOLOGY: linear
 MOLECULE TYPE: protein
 US-08-669-656A-6

Query Match 46.4%; Score 13; DB 4; Length 2132;
 Best Local Similarity 100.0%; Pred. No. 0.00013;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 FVLVIFLGSFYL 16
 DB 375 FVLVIFLGSFYL 387

RESULT 24
 US-09-562-737-83
 Sequence 83, Application US/09562737
 Patent No. 6428967
 GENERAL INFORMATION:
 APPLICANT: Herz, Joachim
 APPLICANT: Goethardt, Michael
 TITLE OF INVENTION: LDL Receptor Signaling Pathways
 FILE REFERENCE: UTSW0708
 CURRENT APPLICATION NUMBER: US/09/562,737
 CURRENT FILING DATE: 2000-05-01
 NUMBER OF SEQ ID NOS: 132
 SOFTWARE: PatentIn Ver. 2.1
 SEQ ID NO 83
 LENGTH: 1024
 TYPE: PRT
 ORGANISM: Artificial Sequence
 FEATURE:
 OTHER INFORMATION: Description of Artificial Sequence: Synthetic
 US-09-562-737-83

Query Match 42.9%; Score 12; DB 4; Length 1024;
 Best Local Similarity 100.0%; Pred. No. 0.00066;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 FVLVIFLGSFYL 16
 DB 404 FVLVIFLGSFYL 415

RESULT 25
 US-09-354-147C-7
 Sequence 7, Application US/09354147C
 Patent No. 6573067
 GENERAL INFORMATION:
 APPLICANT: Dib-Hajj, Sulayman
 APPLICANT: Waxman, Stephen G.
 TITLE OF INVENTION: Modulation of Sodium Channels in Dorsal Root Ganglia
 FILE REFERENCE: 44574-5004-01-US
 CURRENT APPLICATION NUMBER: US/09/354,147C
 CURRENT FILING DATE: 1999-07-16
 PRIOR APPLICATION NUMBER: US 60/072,990
 PRIOR FILING DATE: 1998-01-29
 PRIOR APPLICATION NUMBER: US 60/109,402
 PRIOR FILING DATE: 1998-11-20
 PRIOR APPLICATION NUMBER: PCT/US99/02008
 PRIOR FILING DATE: 1999-01-29
 NUMBER OF SEQ ID NOS: 44
 SOFTWARE: PatentIn Ver. 2.1
 SEQ ID NO 7
 LENGTH: 1233
 TYPE: PRT
 ORGANISM: Homo sapiens
 FEATURE:
 NAME/KEY: UNSURE
 LOCATION: (308)

OTHER INFORMATION: Xaa is leu. Xaa results from a "y" in SEQ ID NO: 6.
 US-09-354-147C-7

Query Match 42.9%; Score 12; DB 4; Length 1233;
 Best Local Similarity 100.0%; Pred. No. 0.00078;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 VIFLGSFYLLNL 19
 DB 184 VIFLGSFYLLNL 195

Search completed: January 27, 2005, 17:54:17
 Job time : 23.5 secs

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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 27, 2005, 17:32:04 ; Search time 86.5 Seconds
(without alignments)
116.120 Million cell updates/sec

Title: US-10-608-584-13
Perfect score: 28
Sequence: 1 CLTVFMVMVIGNLVVLFLALLLSF 28

Scoring table: OLIGO
Gapop 60.0 , Gapext 60.0

Searched: 2002273 seqs, 358729299 residues

Word size : 0

Total number of hits satisfying chosen parameters: 2002273

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Listing first 100 summaries

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- 2: geneseqp1990s:*
- 3: geneseqp2000s:*
- 4: geneseqp2001s:*
- 5: geneseqp2002s:*
- 6: geneseqp2003as:*
- 7: geneseqp2003bs:*
- 8: geneseqp2004s:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	* Query Match	Length	DB ID	Description
1	28	100.0	1795	7	ADB78596 Human sod
2	28	100.0	1855	7	ADB78597 Human sod
3	28	100.0	1981	7	ABR83185 Human SCN
4	28	100.0	1998	7	ABR83184 Human SCN
5	28	100.0	1999	5	ABBO6026 Human adu
6	28	100.0	2005	4	AAB99676 Human adu
7	28	100.0	2005	4	AAB99677 Human neo
8	28	100.0	2005	5	ABBR83627 Human GRF
9	28	100.0	2005	7	ADB78604 Human sod
10	28	100.0	2005	7	ADB78603 Human sod
11	28	100.0	2005	7	ADB78605 Human sod
12	28	100.0	2005	7	ADC46947 Human SCN
13	28	100.0	2009	4	AAB99674 Human adu
14	28	100.0	2009	5	ABG69292 Human sod
15	28	100.0	2009	5	ABG68291 Human sod
16	28	100.0	2009	5	ABG68293 Human sod
17	28	100.0	2009	5	ABG69289 Human sod
18	28	100.0	2009	5	ABG69290 Human sod
19	28	100.0	2009	5	ABBR83626 Human GRF
20	28	100.0	2009	5	AAE16776 Human tra
21	28	100.0	2009	7	ADB78599 Human sod
22	28	100.0	2009	7	ADB78595 Human sod
23	28	100.0	2009	7	ADB78593 Human sod
24	28	100.0	2009	7	ADB78594 Human sod
25	28	100.0	2009	7	ADB78598 Human sod

26	28	100.0	2009	7	ABR83180 Human SCN
27	28	100.0	2009	7	ADE57563 Human Pro
28	28	89.3	1976	7	ADE57386 Rat Prote
29	25	89.3	1978	2	AAW69361 Tetrodoto
30	25	89.3	1980	3	AAW69363 Tetrodoto
31	25	89.3	1980	5	AAO14927 Human sod
32	25	89.3	1980	7	ADB78600 Human sod
33	25	89.3	1980	7	ADB78606 Human sod
34	25	89.3	1988	2	AAW69362 Tetrodoto
35	24	85.7	1381	5	AAE20513 Human ion
36	24	85.7	1382	5	AAE20514 Human ion
37	24	85.7	1392	5	AAE20518 Human ion
38	24	85.7	1398	5	AAE20519 Human ion
39	24	85.7	1442	5	AAE20512 Human ion
40	24	85.7	1453	5	AAE20517 Human ion
41	24	85.7	1962	5	AAE20511 Human ion
42	24	85.7	1973	5	AAE20516 Human ion
43	24	85.7	1998	5	AAE20510 Human ion
44	24	85.7	2009	5	AAE20515 Human ion
45	24	82.1	1835	2	AAE92316 Periphra
46	23	82.1	1977	8	AAE99641 Periphra
47	23	82.1	1977	8	ADE77202 Human pro
48	23	82.1	1977	8	ADL13028 Human ate
49	23	82.1	1978	7	ADE34553 Human Pro
50	23	82.1	1978	7	ADE34549 Human Pro
51	23	82.1	1984	2	AAE99639 Periphra
52	23	82.1	1984	7	ADE54547 Rat Prote
53	23	82.1	1984	7	ADE54551 Rat Prote
54	23	82.1	1984	7	ADE63027 Rat Prote
55	23	82.1	1989	2	AAE92317 Periphra
56	23	82.1	1989	2	AAE99640 Human Pro
57	22	78.6	1836	7	ADE57388 Human Pro
58	22	78.6	1836	7	ADE59630 Human Pro
59	22	78.6	1836	7	ADE63029 Human Pro
60	22	78.6	1836	8	ADQ17412 Human sof
61	21	75.0	1024	5	ABBO4858 LDL recep
62	21	75.0	1107	6	ABR41495 Human DIT
63	21	75.0	1366	7	ADJ68488 Human hea
64	21	75.0	1366	8	ADL06575 Human tum
65	21	75.0	1950	4	ADB78607 Human sod
66	21	75.0	1951	4	ABR99678 Human adu
67	21	75.0	1951	4	ABR99679 Human neo
68	21	75.0	1951	4	ABR99676 Rat Prote
69	21	75.0	1951	8	ADL06576 Human tum
70	21	75.0	2000	5	ABBO6027 Human sod
71	21	75.0	2000	8	ADK81762 Human Nav
72	21	75.0	2015	4	ABR82242 Human SCN
73	21	75.0	2015	7	ADP56441 Human Nav
74	21	75.0	2015	8	ADM34001 Human SCN
75	21	75.0	2015	8	ADM33999 Human SCN
76	21	75.0	2016	4	AAW23994 Human h1
77	21	75.0	2016	4	ABR82239 Human SCN
78	21	75.0	2016	4	ABR82240 Human SCN
79	21	75.0	2016	4	ABR82245 Human SCN
80	21	75.0	2016	4	ABR82243 Human SCN
81	21	75.0	2016	4	ABR82244 Human SCN
82	21	75.0	2016	7	AAE82241 Human SCN
83	21	75.0	2016	7	ADD44756 Human Pro
84	21	75.0	2016	7	ADE55106 Human Pro
85	21	75.0	2016	8	ADM33997 Human SCN
86	21	75.0	2016	8	ADM33995 Human SCN
87	21	75.0	2019	2	AAE67913 Cardiac s
88	17	60.7	2100	2	AAW89579 Cardiac p
89	17	60.7	2105	2	AAW57772 Musca dom
90	17	60.7	2105	2	AAW89577 Calciun p
91	17	60.7	2131	4	ABW64743 Drosophi1
92	17	60.7	2131	8	ADL10141 Drosophi1
93	17	60.7	2131	8	ADL10146 Drosophi1
94	16	57.1	1011	2	AAE99638 Periphra
95	16	57.1	1011	2	AAE57561 Rat Prote
96	13	46.4	115	5	ABG37774 Human pep
97	12	42.9	2104	2	AAW57773 Musca dom
98	12	42.9	2104	2	AAW89578 Calciun p

99 10 35.7 125 8 ABO54760
100 10 35.7 2020 2 AAR06584

ABO54760 Human gen
AAR06584 Cardiac s

ALIGNMENTS

RESULT 1
ADB78596
ID ADB78596 standard; protein, 1795 AA.

AC ADB78596;
XX
XX 04-DEC-2003 (first entry)

DE Human sodium channel subunit mutant SEQ ID NO:140.

KW mutein; mutant; ion channel; ion channel subunit; ICS; nootropic;
KW neuroprotective; ionotropic; antipyrctic; antiarrhythmic; antimigraine;
KW antidepressant; antiparkinsonian; neuroleptic; tranquiliser; analgesic;
KW nephrotoxic; antidiabetic; ophthalmological; epilepsy;
KW ion channel dysfunction; human.

OS Synthetic.
XX Homo sapiens.

XX WO2003008574-A1.

XX 30-JAN-2003.

XX 08-JUL-2002; 2002MO-AU000910.

XX 18-JUL-2001; 2001AU-00006452.

XX 05-MAR-2002; 2002AU-00000910.

XX 13-MAY-2002; 2002AU-00002292.

(BION-) BIONOMICS LTD.
(WALL/) WALLACE R W.

PI Mulley JC, Harkin LA, Dibbens LM, Phillips HA, Heron SE;
PI Berkovic SF, Scheffer IE;

DR MPI; 2003-239332/23.

XX N-PSDB; ADB78635.

PT Identifying predisposition to an ion channel dysfunction, such as
PT periodic paralysis, cardiac arrhythmias, migraine, Alzheimer's disease,
PT schizophrenia, anxiety and depression, by detecting encoding-gene
PT mutation events.

PS Claim 13; SEQ ID NO 140; 106bp; English.

CC The invention relates to a novel method for identifying a subject
CC predisposed to a disorder associated with ion channel dysfunction. The
CC method comprises ascertaining if at least one of the genes encoding ion
CC channel subunits (ICS) has undergone a mutation event so that a cDNA
CC derived from the subject has any of 134 nucleotide sequences. The method
CC of the invention has nootropic, neuroprotective, inotropic, antipyrctic,
CC antiarrhythmic, antimigraine, antidepressant, antiparkinsonian,
CC ophthalmological activity. A polynucleotide of the invention acts as an
CC ion channel agonist, or ion channel antagonist. The methods, isolated
CC nucleic acids, polypeptides, antibody, selective agonist, antagonist or
CC modulator of an ion channel, cells and genetically modified non-human
CC animal, are useful for the diagnosis and treatment of epilepsy and/or a
CC disorder associated with ion channel dysfunction, such as hyper- or hypo-
CC kalemic periodic paralysis, myotonia, malignant hyperthermia, Alz-
CC myasthenia, cardiac arrhythmias, episodic ataxia, migraine, Alzheimer's
CC disease, Parkinson's disease, schizophrenia, hyperplexia, anxiety,
CC depression, phobic obsessive symptoms, neuropathic pain, inflammatory
CC pain, chronic/acute pain, Barter's syndrome, polycystic kidney disease,
CC Dent's disease, hypernatraemic hypoglycaemia of infancy, cystic
CC fibrosis, congenital stationary night blindness and total colour

CC blindness. The present sequence represents a mutant protein of the
CC invention. The sequence data for this patent is not represented in the
CC printed specification, but was obtained in electronic format directly
CC from WIPO at ftp.wipo.int/pub/published_pat_sequences.

XX SQ Sequence 1795 AA;

Query Match 100.0%; Score 28; DB 7; Length 1795;
Best Local Similarity 100.0%; Pred. NO. 6.7e-18;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CLTIVMMVMTGMLVNLPLALLSSP 28
Db 968 CLTIVMMVMTGMLVNLPLALLSSP 995

RESULT 2
ADB78597
ID ADB78597 standard; protein, 1855 AA.

AC ADB78597;

XX 04-DEC-2003 (first entry)

DE Human sodium channel subunit mutant SEQ ID NO:141.

KW mutein; mutant; ion channel; ion channel subunit; ICS; nootropic;
KW neuroprotective; ionotropic; antipyrctic; antiarrhythmic; antimigraine;
KW antidepressant; antiparkinsonian; neuroleptic; tranquiliser; analgesic;
KW nephrotoxic; antidiabetic; ophthalmological; epilepsy;
KW ion channel dysfunction; human.

OS Synthetic.

XX Homo sapiens.

XX WO2003008574-A1.

XX 30-JAN-2003.

XX 08-JUL-2002; 2002MO-AU000910.

XX 18-JUL-2001; 2001AU-00006452.

XX 05-MAR-2002; 2002AU-00000910.

XX 13-MAY-2002; 2002AU-00002292.

(BION-) BIONOMICS LTD.
(WALL/) WALLACE R W.

PI Mulley JC, Harkin LA, Dibbens LM, Phillips HA, Heron SE;
PI Berkovic SF, Scheffer IE;

DR MPI; 2003-239332/23.

XX N-PSDB; ADB78636.

PT Identifying predisposition to an ion channel dysfunction, such as
PT periodic paralysis, cardiac arrhythmias, migraine, Alzheimer's disease,
PT schizophrenia, anxiety and depression, by detecting encoding-gene
PT mutation events.

PS Claim 13; SEQ ID NO 141; 106bp; English.

CC The invention relates to a novel method for identifying a subject
CC predisposed to a disorder associated with ion channel dysfunction. The
CC method comprises ascertaining if at least one of the genes encoding ion
CC channel subunits (ICS) has undergone a mutation event so that a cDNA
CC derived from the subject has any of 134 nucleotide sequences. The method
CC of the invention has nootropic, neuroprotective, inotropic, antipyrctic,
CC antiarrhythmic, antimigraine, antidepressant, antiparkinsonian,
CC neuroleptic, tranquiliser, analgesic, neuroprotective, inotropic, antipyrctic,
CC ophthalmological activity. A polynucleotide of the invention acts as an
CC ion channel agonist, or ion channel antagonist. The methods, isolated
CC nucleic acids, polypeptides, antibody, selective agonist, antagonist or
CC modulator of an ion channel, cells and genetically modified non-human

CC animal, are useful for the diagnosis and treatment of epilepsy and/or a
 CC disorder associated with ion channel dysfunction, such as hyper- or hypo-
 CC kalemic periodic paralysis, myotonias, malignant hyperthermia,
 CC myasthenia, cardiac arrhythmias, episodic ataxia, migraine, Alzheimer's
 CC disease, Parkinson's disease, schizophrenia, hyperkplexia, anxiety,
 CC depression, phobic obsessive symptoms, neuropathic pain, inflammatory
 CC pain, chronic/acute pain, Bartter's syndrome, polycystic kidney disease,
 CC Dent's disease, hyperinsulinaemic hypoglycaemia of infancy, cystic
 CC fibrosis, congenital stationary night blindness and total colour
 CC blindness. The present sequence represents a mutant protein of the
 CC invention. The sequence data for this patent is not represented in the
 CC printed specification, but was obtained in electronic format directly
 CC from WIPO at ftp.wipo.int/pub/published_pat_sequences.

CC Sequence 1855 AA;

Query Match 100.0%; Score 28; DB 7; Length 1855;
 Best Local Similarity 100.0%; Pred. No. 6,9e-18;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 CLTVFMVMVYIGNLVNLFPLALLSSP 28
 DB 968 CLTVFMVMVYIGNLVNLFPLALLSSP 995

RESULT 3

ID ABR83185 standard; protein; 1981 AA.

AC ABR83185;

DT 15-JAN-2004 (first entry)

DE Human SCN1A splice variant -84P:SCN1ADELP654-681.

XX SCN1A; sodium channel type 1 alpha-subunit; anticonvulsant; analgesic;
 KW neuroprotective; anesthetic; cytosolic; cerebroprotective; cardiant;
 KM hypotensive; gene therapy; human; splice variant.

XX Homo sapiens.

XX WO2003072751-A2.

XX 04-SEP-2003.

XX 25-FEB-2003; 2003WO-US006010.

XX 25-FEB-2002; 2002US-0359382P.

XX (UYVA-) UNIV VANDERBILT.

XX George AL, Lossein C;

XX WPI; 2003-712725/67.

XX N-PSDB; ACF57880.

PT Recombinantly expressed sodium channel type 1 alpha subunit, useful in
 screening for modulators, for treating e.g. epilepsy.

XX Disclosure; Page 162-169; 176pp; English.

CC The invention relates to a recombinantly expressed and isolated human
 CC SCN1A (sodium channel type 1 alpha-subunit) (I). (I), optionally
 CC incorporated into a cell, is used to screen for specific modulators,
 CC potentially useful as anticonvulsant, antiepileptic, neuroprotective,
 CC analgesic and/or anesthetic agents, e.g. for treating severe myoclonic
 CC epilepsy of infancy, stroke, cardiac arrest, hyperkalemic paralysis,
 CC motor endplate diseases, hypertension, congestive heart failure and
 CC muscular dystrophy also to treat cancer (SCN1A is expressed in prostatic
 CC and metastatic cancer cell lines). These activities can also be provided
 CC by gene therapy vectors that express (I) or the modulators. The
 CC modulators, also antibodies directed against (I), are used to detect
 CC sodium channel polypeptides. The present sequence represents a human

CC SCN1A splice variant 84P:SCN1ADELP654-681
 XX Sequence 1981 AA;

Query Match 100.0%; Score 28; DB 7; Length 1981;
 Best Local Similarity 100.0%; Pred. No. 7,3e-18;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 CLTVFMVMVYIGNLVNLFPLALLSSP 28
 DB 940 CLTVFMVMVYIGNLVNLFPLALLSSP 967

RESULT 4

ID ABR83184 standard; protein; 1998 AA.

AC ABR83184;

DT 15-JAN-2004 (first entry)

DE Human SCN1A splice variant -33P:SCN1ADELP671-681.

XX SCN1A; sodium channel type 1 alpha-subunit; anticonvulsant; analgesic;
 KW neuroprotective; anesthetic; cytosolic; cerebroprotective; cardiant;
 KM hypotensive; gene therapy; human; splice variant.

XX Homo sapiens.

XX WO2003072751-A2.

XX 04-SEP-2003.

XX 25-FEB-2003; 2003WO-US006010.

XX 25-FEB-2002; 2002US-0359382P.

XX (UYVA-) UNIV VANDERBILT.

XX George AL, Lossein C;

XX WPI; 2003-712725/67.

XX N-PSDB; ACF57879.

PT Recombinantly expressed sodium channel type 1 alpha subunit, useful in
 screening for modulators, for treating e.g. epilepsy.

XX Disclosure; Page 148-156; 176pp; English.

CC The invention relates to a recombinantly expressed and isolated human
 CC SCN1A (sodium channel type 1 alpha-subunit) (I). (I), optionally
 CC incorporated into a cell, is used to screen for specific modulators,
 CC potentially useful as anticonvulsant, antiepileptic, neuroprotective,
 CC analgesic and/or anesthetic agents, e.g. for treating severe myoclonic
 CC epilepsy of infancy, stroke, cardiac arrest, hyperkalemic paralysis,
 CC motor endplate diseases, hypertension, congestive heart failure and
 CC muscular dystrophy also to treat cancer (SCN1A is expressed in prostatic
 CC and metastatic cancer cell lines). These activities can also be provided
 CC by gene therapy vectors that express (I) or the modulators. The
 CC modulators, also antibodies directed against (I), are used to detect
 CC sodium channel polypeptides. The present sequence represents a human
 CC SCN1A splice variant 33P:SCN1ADELP671-681 encoding cDNA

XX Sequence 1998 AA;

Query Match 100.0%; Score 28; DB 7; Length 1998;
 Best Local Similarity 100.0%; Pred. No. 7,4e-18;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 CLTVFMVMVYIGNLVNLFPLALLSSP 28
 DB 957 CLTVFMVMVYIGNLVNLFPLALLSSP 984

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RESULT 5
AB06026
ID ABB06026 standard; protein; 1999 AA.
XX
AC ABB06026;
XX
DT 10-MAY-2002 (first entry)
XX
DE Human sodium channel SCN1A protein SEQ ID NO:2.
XX
KW Human; sodium channel; SCN1A; chromosome 2q24;
KM familial hypercalcaemic periodic paralysis; motor endplate disease.
XX
OS Homo sapiens.
XX
PN WO200196552-A1.
XX
PD 20-DEC-2001.
XX
PF 12-JUN-2001; 2001WO-JP004956.
XX
PR 13-JUN-2000; 2000JP-00177540.
XX
PR 13-JUN-2000; 2000JP-00177544.
XX
PA (NISC-) JAPAN SCI & TECHNOLOGY CORP.
XX
PI Kanazawa I, Goto J, Jeong S;
XX
DR WPI; 2002-098066/13.
XX
DR N-PSDB; ABL39689.
XX
PT Human sodium channels SCN1A and SCN3A and encoded genes, useful in
XX studying physiological mechanism in which excitant cells participate and
XX causes of diseases and developing drugs for motor endplate disease.
XX
PS Claim 1; Page 40-49; 88pp; Japanese.
XX
CC The present invention describes human sodium channels SCN1A and SCN3A.
XX CC SCN3A have been located to the human chromosome 2 long arm, positions
XX CC 2q24 and 2q24-31 respectively. The sodium channel proteins are useful in
XX CC studying the physiological mechanism in which excitant cells participate
XX CC and cause human diseases, and in developing remedies for e.g. familial
XX CC hypercalcaemic periodic paralysis of extremities and motor endplate
XX CC disease
XX
SQ Sequence 1999 AA;
XX
Query Match 100.0%; Score 28; DB 5; Length 1999;
Best Local Similarity 100.0%; Pred. No. 7.4e-18;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTVMVMVIGNVLVNLFLALLLSF 28
DB 958 CTTVMVMVIGNVLVNLFLALLLSF 985

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XX
PN WO200138564-A2.
XX
PD 31-MAY-2001.
XX
PF 24-NOV-2000; 2000WO-CA001404.
XX
PR 26-NOV-1999; 99US-0167623P.
XX
PA (UYMC-) UNIV MCGILL.
XX
PI Rouleau GA, Lafreniere RG, Rochefort D, Coesette P, Ragsdale D;
XX
DR WPI; 2001-355945/37.
XX
DR N-PSDB; AAH55793.
XX
PT Determining a predisposition to epilepsy and/or development of epilepsy
XX PT comprises determining the genotype of SCN1A, SCN2A and/or SCN3A, or a DNA
XX variant, equivalent, or mutation which shows a linkage disequilibrium.
XX
PS Disclosure; Page 123-130; 268pp; English.
XX
CC The present invention describes a method (M1) of determining an
XX CC individual's predisposition to epilepsy and/or development of epilepsy,
XX CC as well as predicting the individual's response to medication. The method
XX CC comprises determining the genotype of at least one gene selected from
XX CC SCN1A, SCN2A or SCN3A, or a DNA variant, equivalent or mutation which
XX CC shows a linkage disequilibrium. SCN1A, SCN2A and SCN3A are all sodium
XX CC channel genes located on chromosome 2. The idiopathic generalised
XX CC epilepsy (IGE) gene is more specifically localised on chromosome 2q23-
XX CC q31. Compounds identified as modulators of the biological activity of
XX CC SCN1A, SCN2A or SCN3A proteins or genes, are useful for treating epilepsy
XX CC or other neurological disorders. They have anticonvulsant and
XX CC neuroprotective activities. AAH55763 to AAH56164 and AAB99674 to AAB99679
XX CC represent SCN1A, SCN2A, and SCN3A cDNAs, gene fragments, PCR primers,
XX CC oligonucleotides and proteins given in the exemplification of the present
XX CC invention
XX
SQ Sequence 2005 AA;
XX
Query Match 100.0%; Score 28; DB 4; Length 2005;
Best Local Similarity 100.0%; Pred. No. 7.4e-18;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 CTTVMVMVIGNVLVNLFLALLLSF 28
DB 959 CTTVMVMVIGNVLVNLFLALLLSF 986

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RESULT 6
AAB99676
ID AAB99676 standard; protein; 2005 AA.
XX
AC AAB99676;
XX
DT 04-SEP-2001 (first entry)
XX
DE Human adult form of SCN2A protein sequence SEQ ID NO:35.
XX
KW Human; epilepsy; chromosome 2; SCN1A; SCN2A; SCN3A; identification;
KM diagnosis; mutation; chromosome 2q23-q31; neurological disorder;
XX anticonvulsant; neuroprotective.
XX
OS Homo sapiens.

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RESULT 7
AAB99677
ID AAB99677 standard; protein; 2005 AA.
XX
AC AAB99677;
XX
DT 04-SEP-2001 (first entry)
XX
DE Human neonatal form of SCN2A protein sequence SEQ ID NO:36.
XX
KW Human; epilepsy; chromosome 2; SCN1A; SCN2A; SCN3A; identification;
KM diagnosis; mutation; chromosome 2q23-q31; neurological disorder;
XX anticonvulsant; neuroprotective.
XX
OS Homo sapiens.
XX
PN WO200138564-A2.
XX
PD 31-MAY-2001.
XX
PF 24-NOV-2000; 2000WO-CA001404.
XX
PR 26-NOV-1999; 99US-0167623P.
XX

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PA (UYMC-) UNIV MCGILL.
 PI Rouleau GA, Lafreniere RG, Rochefort D, Cossette P, Ragsdale D;
 XX WPI; 2001-355945/37.
 DR N-PSDB; AAH55794.
 XX
 PT Determining a predisposition to epilepsy and/or development of epilepsy
 PT comprises determining the genotype of SCN1A, SCN2A and/or SCN3A, or a DNA
 PT variant, equivalent, or mutation which shows a linkage disequilibrium.
 XX
 PS Disclosure; Page 131-138; 268pp; English.
 XX
 CC The present invention describes a method (M1) of determining an
 CC individual's predisposition to epilepsy and/or development of epilepsy,
 CC as well as predicting the individual's response to medication. The method
 CC comprises determining the genotype of at least one gene selected from
 CC SCN1A, SCN2A or SCN3A, or a DNA variant, equivalent, or mutation which
 CC shows a linkage disequilibrium. SCN1A, SCN2A and SCN3A are all sodium
 CC channel genes located on chromosome 2. The idiopathic generalized
 CC epilepsy (IGE) gene is more specifically localised on chromosome 2q23-
 CC q31. Compounds identified as modulators of the biological activity of
 CC SCN1A, SCN2A or SCN3A proteins or genes, are useful for treating epilepsy
 CC or other neurological disorders. They have anticonvulsant and
 CC neuroprotective activities. AAH55763 to AAH56164 and AAH59674 to AAH59679
 CC represent SCN1A, SCN2A, and SCN3A cDNAs, gene fragments, PCR primers,
 CC oligonucleotides and proteins given in the exemplification of the present
 CC invention
 XX
 SQ Sequence 2005 AA;
 XX
 QY Query Match 100.0%; Score 28; DB 4; Length 2005;
 Db Best Local Similarity 100.0%; Pred. No. 7,4e-18; Mismatches 0; Gaps 0;
 Matches 28; Conservative 0; Indels 0; Gaps 0;
 QY 1 CLTVFMVMVIGNLVNLFLALLLSF 28
 Db 959 CLTVFMVMVIGNLVNLFLALLLSF 986
 XX
 RESULT 8
 ABB83627
 ID ABB83627 standard; protein; 2005 AA.
 XX
 AC ABB83627;
 XX
 DT 10-OCT-2002 (first entry)
 XX
 DE Human GEFs+ protein with SCN2A mutation.
 XX
 KW Human; GEFs+; SCN2A; mutant; muten;
 KW generalized epilepsy with febrile seizure plus.
 XX
 OS Homo sapiens.
 OS
 PN JP2002136289-A.
 XX
 PD 14-MAY-2002.
 XX
 PF 01-NOV-2000; 2000JP-00334969.
 XX
 PR 01-NOV-2000; 2000JP-00334969.
 XX
 PA (KAGA-) KAGAKU GIUTSU SHINKO JIGYODAN.
 PA (RIKA) RIKAGAKU KENKYUSHO.
 XX
 DR WPI; 2002-552308/59.
 DR N-PSDB; ABQ79201.
 XX
 PT A human polynucleotide which is complementary to an mRNA transcribed from
 PT a generalized epilepsy with febrile seizure plus (GEFS+)-related gene
 PT useful for diagnosing GEFS+.
 XX

PS Claim 10; Page 29-34; 37pp; Japanese.
 XX
 CC This invention relates to a human polynucleotide which is complementary
 CC to an mRNA transcribed from a "generalized epilepsy with febrile seizure
 CC plus" (GEFS+)-related gene. The gene is useful for diagnosing GEFS+. The
 CC present sequence represents the human GEFs+ protein sequence with SCN2A
 CC mutation
 XX
 SQ Sequence 2005 AA;
 XX
 QY Query Match 100.0%; Score 28; DB 5; Length 2005;
 Db Best Local Similarity 100.0%; Pred. No. 7,4e-18; Mismatches 0; Gaps 0;
 Matches 28; Conservative 0; Indels 0; Gaps 0;
 QY 1 CLTVFMVMVIGNLVNLFLALLLSF 28
 Db 959 CLTVFMVMVIGNLVNLFLALLLSF 986
 XX
 RESULT 9
 ADB78604
 ID ADB78604 standard; protein; 2005 AA.
 XX
 AC ADB78604;
 XX
 DT 04-DEC-2003 (first entry)
 XX
 DE Human sodium channel subunit mutant SEQ ID NO:148.
 XX
 KW muten; mutant; ion channel; ion channel subunit; ICS; noctropic;
 KW neuroprotective; inotropic; antipyretic; antiarrhythmic; antianginal;
 KW antidepressant; antiparkinsonian; neuroleptic; tranquilizer; analgesic;
 KW nephrotoxic; antidiabetic; ophthalmological; epilepsy;
 KW ion channel dysfunction; human.
 XX
 OS Synthetic.
 OS
 PN WO2003008574-A1.
 XX
 PD 30-JAN-2003.
 XX
 PF 08-JUL-2002; 2002WO-AU000910.
 XX
 PR 18-JUL-2001; 2001AU-00006452.
 PR 05-MAR-2002; 2002AU-00000910.
 PR 13-MAY-2002; 2002AU-00002292.
 XX
 PA (BION-) BIONOMICS LTD.
 PA (WALL) WALLACE R W.
 XX
 PI Mulley JC, Harkin LA, Dibbens LM, Phillips HA, Heron SE;
 PI Berkovic SF, Scheffer IE;
 XX
 DR WPI; 2003-239332/23.
 DR N-PSDB; ADB78643.
 XX
 PT Identifying predisposition to an ion channel dysfunction, such as
 PT periodic paralysis, cardiac arrhythmias, migraine, Alzheimer's disease,
 PT schizophrenia, anxiety and depression, by detecting encoding-gene
 PT mutation events.
 XX
 PS Claim 13; SEQ ID NO 148; 106pp; English.
 XX
 CC The invention relates to a novel method for identifying a subject
 CC predisposed to a disorder associated with ion channel dysfunction. The
 CC method comprises ascertaining if at least one of the genes encoding ion
 CC channel subunits (ICS) has undergone a mutation event so that a cDNA
 CC derived from the subject has any of 134 nucleotide sequences. The method
 CC of the invention has noctropic, neuroprotective, inotropic, antipyretic,
 CC antiarrhythmic, antianginal, antidepressant, antiparkinsonian,
 CC neuroleptic, tranquilizer, analgesic, nephrotoxic, antidiabetic, and
 CC ophthalmological activity. A polynucleotide of the invention acts as an

CC ion channel agonist, or ion channel antagonist. The methods, isolated
CC nucleic acids, polypeptides, antibody, selective agonist, antagonist or
CC modulator of an ion channel, cells and genetically modified non-human
CC animal, are useful for the diagnosis and treatment of epilepsy and/or a
CC disorder associated with ion channel dysfunction, such as hyper- or hypo-
CC kalemia, periodic paralysis, myotonia, malignant hyperthermia,
CC myasthenia, cardiac arrhythmias, episodic ataxia, migraine, Alzheimer's
CC disease, Parkinson's disease, schizophrenia, hyperkplexia, anxiety,
CC depression, phobic obsessive symptoms, neuropathic pain, inflammatory
CC pain, chronic/acute pain, Bartter's syndrome, polycystic kidney disease,
CC Dent's disease, hyperinsulinaemic hypoglycaemia of infancy, cystic
CC fibrosis, congenital stationary night blindness and total colour
CC blindness. The present sequence represents a mutant protein of the
CC invention. The sequence data for this patent is not represented in the
CC printed specification, but was obtained in electronic format directly
CC from WIPO at ftp.wipo.int/pub/published_pat_sequences.

SQ Sequence 2005 AA;

Query Match 100.0%; Score 28; DB 7; Length 2005;
Best Local Similarity 100.0%; Pred. No. 7.4e-18;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CLTFPMWVVGIVLVLNLFLLILSSSF 28
|||
Db 959 CLTFPMWVVGIVLVLNLFLLILSSSF 986

RESULT 10

ADB78603
ID ADB78603 standard; protein; 2005 AA.

XX ADB78603;

DT 04-DEC-2003 (first entry)

DE Human sodium channel subunit mutant SEQ ID NO:147.

XX mutein; mutant; ion channel; ion channel subunit; ICS; nootropic;
KW neuroprotective; inotropic; antipyretic; antiarrhythmic; antimigraine;
KW antidepressant; antiparkinsonian; neuroleptic; tranquiliser; analgesic;
KW nephrotropic; antidiabetic; ophthalmological; epilepsy;
XX ion channel dysfunction; human.

XX Synthetic.

OS Homo sapiens.

XX WO2003008574-A1.

XX 30-JAN-2003.

XX 08-JUL-2002; 2002WO-AU000910.

XX 18-JUL-2001; 2001AU-00006452.

XX 05-MAR-2002; 2002AU-00000910.

XX 13-MAY-2002; 2002AU-00002292.

XX (BION-) BIONOMICS LTD.

XX (WALL/) WALLACE R W.

XX Mulley JC, Harkin LA, Dibbens LM, Phillips HA, Heron SE;

XX Berkovic SF, Scheffer IE;

XX WPI; 2003-239332/23.

XX N-PSDB; ADB78642.

PT Identifying predisposition to an ion channel dysfunction, such as

PT periodic paralysis, cardiac arrhythmias, migraine, Alzheimer's disease,

PT schizophrenia, anxiety and depression, by detecting encoding-gene

PT mutation events.

PS Claim 13; SEQ ID NO 147; 106pp; English.

CC The invention relates to a novel method for identifying a subject
CC predisposed to a disorder associated with ion channel dysfunction. The
CC method comprises ascertaining if at least one of the genes encoding ion
CC channel subunits (ICS) has undergone a mutation event so that a cDNA
CC derived from the subject has any of 134 nucleotide sequences. The method
CC of the invention has nootropic, neuroprotective, inotropic, antipyretic,
CC antiarrhythmic, antimigraine, antidepressant, antiparkinsonian,
CC neuroleptic, tranquiliser, analgesic, nephrotropic, antidiabetic, and
CC ophthalmological activity. A polynucleotide of the invention acts as an
CC ion channel agonist, or ion channel antagonist. The methods, isolated
CC nucleic acids, polypeptides, antibody, selective agonist, antagonist or
CC modulator of an ion channel, cells and genetically modified non-human
CC animal, are useful for the diagnosis and treatment of epilepsy and/or a
CC disorder associated with ion channel dysfunction, such as hyper- or hypo-
CC kalemia, periodic paralysis, myotonia, malignant hyperthermia,
CC myasthenia, cardiac arrhythmias, episodic ataxia, migraine, Alzheimer's
CC disease, Parkinson's disease, schizophrenia, hyperkplexia, anxiety,
CC depression, phobic obsessive symptoms, neuropathic pain, inflammatory
CC pain, chronic/acute pain, Bartter's syndrome, polycystic kidney disease,
CC Dent's disease, hyperinsulinaemic hypoglycaemia of infancy, cystic
CC fibrosis, congenital stationary night blindness and total colour
CC blindness. The present sequence represents a mutant protein of the
CC invention. The sequence data for this patent is not represented in the
CC printed specification, but was obtained in electronic format directly
CC from WIPO at ftp.wipo.int/pub/published_pat_sequences.

SQ Sequence 2005 AA;

Query Match 100.0%; Score 28; DB 7; Length 2005;
Best Local Similarity 100.0%; Pred. No. 7.4e-18;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CLTFPMWVVGIVLVLNLFLLILSSSF 28
|||
Db 959 CLTFPMWVVGIVLVLNLFLLILSSSF 986

RESULT 11

ADB78605
ID ADB78605 standard; protein; 2005 AA.

XX ADB78605;

DT 04-DEC-2003 (first entry)

DE Human sodium channel subunit mutant SEQ ID NO:149.

XX mutein; mutant; ion channel; ion channel subunit; ICS; nootropic;
KW neuroprotective; inotropic; antipyretic; antiarrhythmic; antimigraine;
KW antidepressant; antiparkinsonian; neuroleptic; tranquiliser; analgesic;
KW nephrotropic; antidiabetic; ophthalmological; epilepsy;
XX ion channel dysfunction; human.

XX Synthetic.

OS Homo sapiens.

XX WO2003008574-A1.

XX 30-JAN-2003.

XX 08-JUL-2002; 2002WO-AU000910.

XX 18-JUL-2001; 2001AU-00006452.

XX 05-MAR-2002; 2002AU-00000910.

XX 13-MAY-2002; 2002AU-00002292.

XX (BION-) BIONOMICS LTD.

XX (WALL/) WALLACE R W.

XX Mulley JC, Harkin LA, Dibbens LM, Phillips HA, Heron SE;

XX Berkovic SF, Scheffer IE;

XX WPI; 2003-239332/23.

DR N-PSDB; ADB78644.

XX Identifying predisposition to an ion channel dysfunction, such as
PT periodic paralysis, cardiac arrhythmias, migraine, Alzheimer's disease,
PT schizophrenia, anxiety and depression, by detecting encoding-gene
PT mutation events.

PS Claim 13; SEQ ID NO 149; 106pp; English.

XX The invention relates to a novel method for identifying a subject
XX predisposed to a disorder associated with ion channel dysfunction. The
CC method comprises ascertaining if at least one of the genes encoding ion
CC channel subunits (ICs) has undergone a mutation event so that a cDNA
CC derived from the subject has any of 134 nucleotide sequences. The method
CC of the invention has nootropic, neuroprotective, inotropic, antipyretic,
CC antihypertensive, antimigraine, antidepressant, antiparkinsonian,
CC neuroleptic, tranquilizer, analgesic, nephrotoxic, antidiabetic, and
CC ophthalmological activity. A polynucleotide of the invention acts as an
CC ion channel agonist, or ion channel antagonist. The method, isolated
CC nucleic acids, polypeptides, antibody, selective agonist, antagonist or
CC modulator of an ion channel, cells and genetically modified non-human
CC animal, are useful for the diagnosis and treatment of epilepsy and/or a
CC disorder associated with ion channel dysfunction, such as hyper- or hypo-
CC Kalemic periodic paralysis, myotonia, malignant hyperthermia,
CC myasthenia, cardiac arrhythmias, episodic ataxia, migraine, Alzheimer's
CC disease, Parkinson's disease, schizophrenia, hyperkplexia, anxiety,
CC depression, phobic obsessive symptoms, neuropathic pain, inflammatory
CC pain, chronic/acute pain, Bartter's syndrome, polycystic kidney disease,
CC Dent's disease, hyperinsulinemic hypoglycemia of infancy, cystic
CC fibrosis, congenital stationary night blindness and total colour
CC blindness. The present sequence represents a mutant protein of the
CC invention. The sequence data for this patent is not represented in the
CC printed specification, but was obtained in electronic format directly
CC from WIPO at ftp.wipo.int/pub/published_pat_sequences.

XX Sequence 2005 AA;

Query Match 100.0%; Score 28; DB 7; Length 2005;
Best Local Similarity 100.0%; Pred. No. 7,4e-18;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CLTFFMMWVWVIGNLVNLFLALLSSP 28
Db 959 CLTFFMMWVWVIGNLVNLFLALLSSP 986

RESULT 12

ID ADC46947 standard; protein; 2005 AA.

XX ADC46947;

DT 18-DEC-2003 (first entry)

DE Human SCN2A amino acid sequence #SEQ ID 3.

XX SCN2A; voltage-gated ion channel, human; neuroprotective; gene therapy;
KM vaccine; Alzheimer's disease.

XX Homo sapiens.

PN MO2003060525-A1.

PD 24-JUL-2003.

PF 16-JAN-2003; 2003WO-EP000400.

PR 17-JAN-2002; 2002EP-00001236.

PR 17-JAN-2002; 2002US-0348674P.

PA (EVOT-) EVOTEC NEUROSCIENCES GMBH.

PI Hipfel R, Von Der Kammer H, Pohlner J;

XX WPI; 2003-598580/56.

DR N-PSDB; ADC46947.

XX Diagnosing or prognosticating a neurodegenerative disease by detecting
PT the level or activity of transcription or translation products of the
PT gene coding for the voltage-gated ion channel SCN2A.

PS Disclosure; Fig 9; 67pp; English.

XX The invention relates to a method for diagnosing or prognosticating a
CC neurodegenerative disease in a subject, or determining whether a subject
CC is at increased risk of developing the disease. The method comprises
CC detecting the level and/or activity of a transcription or translation
CC product of the gene coding for the voltage-gated ion channel SCN2A. The
CC modulator of an activity and/or of a level of at least one substance is
CC useful for preparing a composition for treating or preventing a
CC neurodegenerative disease, in particular Alzheimer's disease. The current
CC sequence represents the human SCN2A amino acid sequence.

XX Sequence 2005 AA;

Query Match 100.0%; Score 28; DB 7; Length 2005;
Best Local Similarity 100.0%; Pred. No. 7,4e-18;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CLTFFMMWVWVIGNLVNLFLALLSSP 28
Db 959 CLTFFMMWVWVIGNLVNLFLALLSSP 986

RESULT 13

ID AAB99674 standard; protein; 2009 AA.

XX AAB99674;

DT 04-SEP-2001 (first entry)

DE Human adult form of SCN1A protein sequence SEQ ID NO:3.

XX Human; epilepsy; chromosome 2; SCN1A; SCN2A; SCN3A; identification;
KM diagnosis; mutation; chromosome 2q23-q31; neurological disorder;
XX anticonvulsant; neuroprotective.

XX Homo sapiens.

PN MO200138564-A2.

PD 31-MAY-2001.

PF 24-NOV-2000; 2000WO-CA001404.

PR 26-NOV-1999; 99US-0167623P.

PA (UMC-) UNIV MCGILL.

PT Rouleau GA, Lafreniere RG, Rochefort D, Cossette P, Ragsdale D;

DR WPI; 2001-355945/37.

DR N-PSDB; AAH55763.

XX Determining a predisposition to epilepsy and/or development of epilepsy
PT comprises determining the genotype of SCN1A, SCN2A and/or SCN3A, or a DNA
PT variant, equivalent, or mutation which shows a linkage disequilibrium.

PS Disclosure; Page 96-104; 268pp; English.

XX The present invention describes a method (M1) of determining an
CC individual's predisposition to epilepsy and/or development of epilepsy,
CC as well as predicting the individual's response to medication. The method
CC comprises determining the genotype of at least one gene selected from
CC SCN1A, SCN2A or SCN3A, or a DNA variant, equivalent, or mutation which

CC shows a linkage disequilibrium. SCN1A, SCN2A and SCN3A are all sodium
 CC channel genes located on chromosome 2. The idiopathic generalised
 CC epilepsy (IGE) gene is more specifically localised on chromosome 2q23-
 CC q31. Compounds identified as modulators of the biological activity of
 CC SCN1A, SCN2A or SCN3A proteins or genes, are useful for treating epilepsy
 CC or other neurological disorders. They have anticonvulsant and
 CC neuroprotective activities. AAH55763 to AAH56164 and AAB99674 to AAB99679
 CC represent SCN1A, SCN2A, and SCN3A cDNAs, gene fragments, PCR primers,
 CC oligonucleotides and proteins given in the exemplification of the present
 CC invention
 CC XX
 SQ Sequence 2009 AA;

Query Match 100.0%; Score 28; DB 4; Length 2009;
 Best Local Similarity 100.0%; Pred. No. 7.5e-18;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CLTFFMMVWVIGNLVNLVNLFLALLSSSF 28
 |||
 DB 968 CLTFFMMVWVIGNLVNLVNLFLALLSSSF 995

RESULT 14

ABG69292
 ID ABG69292 standard; protein; 2009 AA.

AC ABG69292;

DT 21-OCT-2002 (first entry)

DE Human sodium channel alpha 1-subunit (SCN1A) variant protein #4.

XX Human; sodium channel alpha 1-subunit; SCN1A; episodic ataxia; epilepsy;
 KW generalised epilepsy with febrile seizures plus; myasthenia;
 KW sodium channel dysfunction; malignant hyperthermia; neuropathic pain;
 KW inflammatory pain; Alzheimer's disease; Parkinson's disease; myotonia;
 KW schizophrenia; hyperkplexia; anticonvulsant; analgesic; neuroprotective;
 KW nocrotropic; anti-Parkinsonian; neuroleptic.

OS Homo sapiens.

XX WO200250096-A1.

PD 27-JUN-2002.

PF 20-DEC-2001; 2001WO-AU001648.

PR 20-DEC-2000; 2000AU-00002203.

PA (BION-) BIONOMICS LTD.

PI Wallace RH, Mulley JC, Berkovic SF;

DR WPI; 2002-528445/56.

XX New nucleic acid encoding mutant alpha subunit of a mammalian voltage-
 PT gated sodium channel, useful for diagnosis of epilepsy, particularly
 PT generalized epilepsy with febrile seizures plus.
 PS Claim 53; Page 147-157; 198pp; English.

XX The invention relates to a nucleic acid molecule encoding a mutant alpha
 CC subunit of a mammalian voltage-gated sodium channel. The DNA and the
 CC polypeptide may be used in the diagnosis of epilepsy, in particular
 CC generalised epilepsy with febrile seizures plus, and other disorders
 CC associated with sodium channel dysfunction. The polypeptide is useful for
 CC the screening of candidate pharmaceutical agents, where high throughput
 CC screening techniques are employed. The sequences are also useful in the
 CC manufacture of a medicament for the treatment of a disorder associated
 CC with sodium channel dysfunction such as epilepsy, particularly
 CC generalised epilepsy with febrile seizures plus, malignant hyperthermia,
 CC myasthenia, episodic ataxia, neuropathic and inflammatory pain,
 CC Alzheimer's disease, Parkinson's disease, schizophrenia, hyperkplexia

CC and myotonia. Sequences ABG69289-ABG69293 represent human sodium channel
 CC alpha 1-subunit (SCN1A) polypeptides of the invention
 CC XX
 SQ Sequence 2009 AA;

Query Match 100.0%; Score 28; DB 5; Length 2009;
 Best Local Similarity 100.0%; Pred. No. 7.5e-18;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CLTFFMMVWVIGNLVNLVNLFLALLSSSF 28
 |||
 DB 968 CLTFFMMVWVIGNLVNLVNLFLALLSSSF 995

RESULT 15

ABG69291
 ID ABG69291 standard; protein; 2009 AA.

AC ABG69291;

DT 21-OCT-2002 (first entry)

DE Human sodium channel alpha 1-subunit (SCN1A) variant protein #3.

XX Human; sodium channel alpha 1-subunit; SCN1A; episodic ataxia; epilepsy;
 KW generalised epilepsy with febrile seizures plus; myasthenia;
 KW sodium channel dysfunction; malignant hyperthermia; neuropathic pain;
 KW inflammatory pain; Alzheimer's disease; Parkinson's disease; myotonia;
 KW schizophrenia; hyperkplexia; anticonvulsant; analgesic; neuroprotective;
 KW nocrotropic; anti-Parkinsonian; neuroleptic.

OS Homo sapiens.

XX WO200250096-A1.

PD 27-JUN-2002.

PF 20-DEC-2001; 2001WO-AU001648.

PR 20-DEC-2000; 2000AU-00002203.

PA (BION-) BIONOMICS LTD.

PI Wallace RH, Mulley JC, Berkovic SF;

DR WPI; 2002-528445/56.

DR N-PADB; ABK98843.

XX New nucleic acid encoding mutant alpha subunit of a mammalian voltage-
 PT gated sodium channel, useful for diagnosis of epilepsy, particularly
 PT generalized epilepsy with febrile seizures plus.
 PS Claim 47; Page 107-117; 198pp; English.

XX The invention relates to a nucleic acid molecule encoding a mutant alpha
 CC subunit of a mammalian voltage-gated sodium channel. The DNA and the
 CC polypeptide may be used in the diagnosis of epilepsy, in particular
 CC generalised epilepsy with febrile seizures plus, and other disorders
 CC associated with sodium channel dysfunction. The polypeptide is useful for
 CC the screening of candidate pharmaceutical agents, where high throughput
 CC screening techniques are employed. The sequences are also useful in the
 CC manufacture of a medicament for the treatment of a disorder associated
 CC with sodium channel dysfunction such as epilepsy, particularly
 CC generalised epilepsy with febrile seizures plus, malignant hyperthermia,
 CC myasthenia, episodic ataxia, neuropathic and inflammatory pain,
 CC Alzheimer's disease, Parkinson's disease, schizophrenia, hyperkplexia
 CC and myotonia. Sequences ABG69289-ABG69293 represent human sodium channel
 CC alpha 1-subunit (SCN1A) polypeptides of the invention
 CC XX

SQ Sequence 2009 AA;

Query Match 100.0%; Score 28; DB 5; Length 2009;
 Best Local Similarity 100.0%; Pred. No. 7.5e-18;

FN	JP2002136289-A.
XX	
PD	14-MAY-2002.
XX	
PF	01-NOV-2000; 2000JP-00334969.
XX	
PR	01-NOV-2000; 2000JP-00334969.
XX	
PA	(KAGA-) KAGAKU GIUTSU SHINKO JIGYODAN. (RIKA) RIKAGAKU KENKYUSHO.
DR	WPI ; 2002-552308/59.
DR	N-PSTDB; ABQ79200.
PT	A human polynucleotide which is complementary to an mRNA transcribed from a generalized epilepsy with febrile seizure plus (GEFS+)-related gene useful for diagnosing GEFS+.
PS	Claim 11; Page 16-21; 37pp; Japanese.
XX	
CC	This invention relates to a human polynucleotide which is complementary to an mRNA transcribed from a "generalized epilepsy with febrile seizure plus" (GEFS+)-related gene. The gene is useful for diagnosing GEFS+. The present sequence represents the human GEFS+ protein sequence with SCN1A mutation
CC	
XX	
SQ	Sequence 2009 AA;
	Query Match 100.0%; Score 28; DB 5; Length 2009; Best Local Similarity 100.0%; Pred. No. 7.5e-18; Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0
OY	1 CLTFFMMVMVIGNLVINLFTALLISSE 28 Db 968 CLTFFMMVMVIGNLVINLFTALLISSE 995
RESULT 20	
AAE16776	
ID	AAE16776 standard; protein; 2009 AA.
XX	
AC	AAE16776;
XX	
DT	09-APR-2002 (first entry)
DE	
XX	
Human transporter and ion channel-13 (TRICH-13) protein.	
KW	Human; transporter and ion channel-13; TRICH-13; neuroprotective; asthma; nootropic; cytosolic; cardiovascular; immunosuppressive; cardiomyopathy; antiinflammatory; protein therapy; akinesia; cystic fibrosis; leukemia; Bell's palsy; amyotrophic lateral sclerosis; Alzheimer's disease; cancer; amaesia; dementia; myocarditis; Duchenne's muscular dystrophy; AIDS; Acquired Immune Deficiency Syndrome; Addison's disease; allergy; angina; cell proliferative disorder; psoriasis; cardiac disease; hypertension; bradyarrhythmia; gene expression; drug screening.
KM	
XX	
OS	Homo sapiens.
XX	
XX	
Key	Location/Qualifiers
FT	Domain 122..147
FT	/label=Transmembrane_domain
FT	Domain 124..422
FT	/note="Ion transport protein domain"
FT	Domain 213..230
FT	/label=Transmembrane_domain
FT	Domain 250..268
FT	/label=Transmembrane_domain
FT	Domain 399..422
FT	/label=Transmembrane_domain
FT	Domain 761..781
FT	/label=Transmembrane_domain
FT	Domain 764..991
FT	/note="Ion transport protein domain"

FT Domain 803..821
 FT /label= Transmembrane_domain
 FT Domain 832..852
 FT /label= Transmembrane_domain
 FT Domain 893..911
 FT /label= Transmembrane_domain
 FT Binding-site 908..915
 FT /label= P loop
 FT /note= "ATP/GTP binding site"
 FT Domain 971..991
 FT /label= Transmembrane_domain
 FT Domain 1214..1482
 FT /note= "Ion transport protein domain"
 FT Domain 1251..1274
 FT /label= Transmembrane_domain
 FT Domain 1350..1369
 FT /label= Transmembrane_domain
 FT Domain 1459..1482
 FT /label= Transmembrane_domain
 FT Domain 1537..1785
 FT /note= "Ion transport protein domain"
 FT Domain 1543..1562
 FT /label= Transmembrane_domain
 FT Domain 1576..1594
 FT /label= Transmembrane_domain
 FT Domain 1602..1620
 FT /label= Transmembrane_domain
 FT Domain 1633..1650
 FT /label= Transmembrane_domain
 FT Domain 1673..1692
 FT /label= Transmembrane_domain
 FT Domain 1762..1785
 FT /label= Transmembrane_domain
 FT Binding-site 1916..1936
 FT /note= "IQ calmodulin-binding motif"
 FT WO200192304-A2.
 PN 06-DEC-2001.
 XX 06-DEC-2001.
 PD 25-MAY-2001; 2001WO-US017065.
 PF 26-MAY-2000; 2000US-0208424P.
 PR 01-JUN-2000; 2000US-0209001P.
 PR 08-JUN-2000; 2000US-0210588P.
 PR 16-JUN-2000; 2000US-0212335P.
 PR 22-JUN-2000; 2000US-0213747P.
 PR 29-JUN-2000; 2000US-0215391P.
 XX (INCY-) INCYTE GENOMICS INC.
 PA Thornton M, Walla NK, Yue H, Nguyen DB, Lai P, Gandhi AR;
 PI Tribouley CM, Yao MG, Rakkumar J, Au-Young J, Lu Y, Tang YF;
 PI Azimzai Y, Bruns CM, Griffin JA, Yang J, Sanjanwala MS, Raumann BE;
 PI Lee RA, Hafalia A, Baughn MR, Green BD, Khan FA, Kearney L;
 PI Ellicot VS, Seilhamer JJ, Policky JL, Borowsky ML, Burford N, Ding L;
 PI Lu DM, Hillman JL;
 XX WPI; 2002-122055/16.
 DR N-PSDB; AAD27266.
 XX New human transporters and ion channels (TRICH) polypeptides useful for
 PT diagnosing, treating or preventing disorders associated with aberrant
 PT expression of TRICH.
 XX Claim 1; Page 158-162; 210pp; English.
 PS The invention relates to human transporters and ion channels (TRICH)
 CC polypeptides and their cDNA molecules. The nucleic acid and polypeptide
 CC sequences are useful in the diagnosis, treatment, and prevention of
 CC disorders associated with transport (akinesia, cystic fibrosis, Bell's
 CC palsy, amyotrophic lateral sclerosis); neurological (Alzheimer's disease,
 CC amnesia, dementia); muscle (cardiomyopathy, myocarditis, Duchenne's

CC muscular dystrophy); immunological (AIDS, Addison's disease, allergies,
 CC asthma); cell proliferative disorders (cancers, leukaemia, psoriasis);
 CC cardiac disease (angina, hypertension, or bradyarrhythmia) and in the
 CC assessment of the effects of exogenous compounds on the expression of
 CC nucleic acid and amino acid sequences of transporters and ion channels.
 CC The polynucleotides may be used to detect and quantify gene expression in
 CC biopsied tissues in which TRICH expression may be correlated with a
 CC disease, to generate hybridization probes for mapping naturally occurring
 CC genomic sequence, and in drug screening. The present sequence is human
 CC TRICH-13 protein
 XX SQ Sequence 2009 AA;
 XX
 XX Query Match 100.0%; Score 28; DB 5; Length 2009;
 XX Best Local Similarity 100.0%; Pred. No. 7.5e-18;
 XX Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 CLTVFMVMVIGNLVNLPLALLISF 28
 Db 968 CLTVFMVMVIGNLVNLPLALLISF 995
 RESULT 21
 ADB78599
 ID ADB78599 standard; protein; 2009 AA.
 XX ADB78599;
 AC ADB78599;
 DT 04-DEC-2003 (first entry)
 XX DE Human sodium channel subunit mutant SEQ ID NO:143.
 XX KM mutein; mutant; ion channel; ion channel subunit; ICS; nootropic;
 KM neuroprotective; inotropic; antipyretic; antiarrhythmic; antimigraine;
 KM antidepressant; antiparkinsonian; neuroleptic; tranquiliser; analgesic;
 KM nephrotoxic; antidiabetic; ophthalmological; epilepsy;
 KM ion channel dysfunction; human.
 XX OS Synthetic.
 OS Homo sapiens.
 XX WO2003008574-A1.
 PN 30-JAN-2003.
 PD 08-JUL-2002; 2002WO-AU000910.
 PF 18-JUL-2001; 2001AU-00006452.
 PR 05-MAR-2002; 2002AU-00000910.
 PR 13-MAY-2002; 2002AU-00002292.
 XX (BION-) BIONOMICS LTD.
 PA (WALL) WALLACE R W.
 XX Mulley JC, Harkin LA, Dibbens LM, Phillips HA, Heron SE;
 PI Berkovic SF, Scheffer IE;
 XX WPI; 2003-239332/23.
 DR N-PSDB; ADB78638.
 XX Identifying predisposition to an ion channel dysfunction, such as
 PT periodic paralysis, cardiac arrhythmias, migraine, Alzheimer's disease,
 PT schizophrenia, anxiety and depression, by detecting encoding-gene
 PT mutation events.
 XX Claim 13; SEQ ID NO 143; 106pp; English.
 PS The invention relates to a novel method for identifying a subject
 CC predisposed to a disorder associated with ion channel dysfunction. The
 CC method comprises ascertaining if at least one of the genes encoding ion
 CC channel subunits (ICS) has undergone a mutation event so that a cDNA
 CC derived from the subject has any of 134 nucleotide sequences. The method
 CC of the invention has nootropic, neuroprotective, inotropic, antipyretic,

CC antiarrhythmic, antimigraine, antidepressant, antiparkinsonian,
CC neuroleptic, tranquilizer, analgesic, neurotropic, antidiabetic, and
CC optunolectic activity. A polynucleotide of the invention acts as an
CC ion channel agonist, or ion channel antagonist. The methods, isolated
CC nucleic acids, polypeptides, antibody, selective agonist, antagonist or
CC modulator of an ion channel, cells and genetically modified non-human
CC animal, are useful for the diagnosis and treatment of epilepsy and/or a
CC disorder associated with ion channel dysfunction, such as hyper- or hypo-
CC kalemic periodic paralysis, myotonia, malignant hyperthermia,
CC myasthenia, cardiac arrhythmias, episodic ataxia, migraine, Alzheimer's
CC disease, Parkinson's disease, schizophrenia, hyperkplexia, anxiety,
CC depression, phobic obsessive symptoms, neuropathic pain, inflammatory
CC pain, chronic/acute pain, Bartter's syndrome, polycystic kidney disease,
CC fibrosis, congenital stationary night blindness and total colour
CC blindness. The present sequence represents a mutant protein of the
CC invention. The sequence data for this patent is not represented in the
CC printed specification, but was obtained in electronic format directly
CC from WIPO at ftp.wipo.int/pub/published_pat_sequences.

XX Sequence 2009 AA;

Query Match 100.0%; Score 28; DB 7; Length 2009;
Best Local Similarity 100.0%; Pred. No. 7.5e-18;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CLTFFMMVWVIGNLVVLNLFALLLSF 28
DB 968 CLTFFMMVWVIGNLVVLNLFALLLSF 995

RESULT 22

ADB78595
ADB78595 standard; protein; 2009 AA.

AC ADB78595;

DT 04-DEC-2003 (first entry)

DE Human sodium channel subunit mutant SEQ ID NO:139.

XX mutant; ion channel; ion channel subunit; ICS; nootropic;
XX neuroprotective; inotropic; antipyretic; antiarrhythmic; antimigraine;
XX antidepressant; antiparkinsonian; neuroleptic; tranquilizer; analgesic;
XX neurotropic; antidiabetic; ophthalmological; epilepsy;
XX ion channel dysfunction; human.

OS Synthetic.

OS Homo sapiens.

PN WO2003008574-A1.

PD 30-JAN-2003.

PE 08-JUL-2002; 2002WO-AU000910.

PR 18-JUL-2001; 2001AU-00006452.

PR 05-MAR-2002; 2002AU-0000910.

PR 13-MAY-2002; 2002AU-00002292.

PA (BION-) BIONOMICS LTD.

PA (WALL/) WALLACE R W.

PI Mulley JC, Harkin LA, Dibbens LM, Phillips HA, Heron SE;

PI Berkovic SF, Scheffer IE;

PI WPI: 2003-239332/23.

PI N-PSDB; ADB78634.

PT Identifying predisposition to an ion channel dysfunction, such as
PT periodic paralysis, cardiac arrhythmias, migraine, Alzheimer's disease,
PT schizophrenia, anxiety and depression, by detecting encoding-gene
PT mutation events.

XX Claim 13; SEQ ID NO 139; 106bp; English.

CC The invention relates to a novel method for identifying a subject
CC predisposed to a disorder associated with ion channel dysfunction. The
CC method comprises ascertaining if at least one of the genes encoding ion
CC channel subunits (ICS) has undergone a mutation event so that a cDNA
CC derived from the subject has any of 134 nucleotide sequences. The method
CC of the invention has nootropic, neuroprotective, inotropic, antipyretic,
CC antiarrhythmic, antimigraine, analgesic, neurotropic, antiparkinsonian,
CC neuroleptic, tranquilizer, analgesic, neurotropic, antidiabetic,
CC ophthalmological activity. A polynucleotide of the invention acts as an
CC ion channel agonist, or ion channel antagonist. The methods, isolated
CC nucleic acids, polypeptides, antibody, selective agonist, antagonist or
CC modulator of an ion channel, cells and genetically modified non-human
CC animal, are useful for the diagnosis and treatment of epilepsy and/or a
CC disorder associated with ion channel dysfunction, such as hyper- or hypo-
CC kalemic periodic paralysis, myotonia, malignant hyperthermia,
CC myasthenia, cardiac arrhythmias, episodic ataxia, migraine, Alzheimer's
CC disease, Parkinson's disease, schizophrenia, hyperkplexia, anxiety,
CC depression, phobic obsessive symptoms, neuropathic pain, inflammatory
CC pain, chronic/acute pain, Bartter's syndrome, polycystic kidney disease,
CC Dent's disease, hyperinsulinaemic hypoglycaemia of infancy, cystic
CC fibrosis, congenital stationary night blindness and total colour
CC blindness. The present sequence represents a mutant protein of the
CC invention. The sequence data for this patent is not represented in the
CC printed specification, but was obtained in electronic format directly
CC from WIPO at ftp.wipo.int/pub/published_pat_sequences.

XX Sequence 2009 AA;

Query Match 100.0%; Score 28; DB 7; Length 2009;
Best Local Similarity 100.0%; Pred. No. 7.5e-18;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CLTFFMMVWVIGNLVVLNLFALLLSF 28
DB 968 CLTFFMMVWVIGNLVVLNLFALLLSF 995

RESULT 23

ADB78593
ADB78593 standard; protein; 2009 AA.

AC ADB78593;

DT 04-DEC-2003 (first entry)

DE Human sodium channel subunit mutant SEQ ID NO:137.

XX mutant; ion channel; ion channel subunit; ICS; nootropic;
XX neuroprotective; inotropic; antipyretic; antiarrhythmic; antimigraine;
XX antidepressant; antiparkinsonian; neuroleptic; tranquilizer; analgesic;
XX neurotropic; antidiabetic; ophthalmological; epilepsy;
XX ion channel dysfunction; human.

OS Synthetic.

OS Homo sapiens.

PN WO2003008574-A1.

PD 30-JAN-2003.

PE 08-JUL-2002; 2002WO-AU000910.

PR 18-JUL-2001; 2001AU-00006452.

PR 05-MAR-2002; 2002AU-0000910.

PR 13-MAY-2002; 2002AU-00002292.

PA (BION-) BIONOMICS LTD.

PA (WALL/) WALLACE R W.

PI Mulley JC, Harkin LA, Dibbens LM, Phillips HA, Heron SE;

PI Berkovic SF, Scheffer IE;
 XX
 DR WPI: 2003-239332/23.
 DR N-PSDB; ADB78632.
 PT Identifying predisposition to an ion channel dysfunction, such as
 PT periodic paralysis, cardiac arrhythmias, migraine, Alzheimer's disease,
 PT schizophrenia, anxiety and depression, by detecting encoding-gene
 PT mutation events.
 XX
 PS Claim 13; SEQ ID NO 137; 106pp; English.
 XX
 CC The invention relates to a novel method for identifying a subject
 CC predisposed to a disorder associated with ion channel dysfunction. The
 CC method comprises ascertaining if at least one of the genes encoding ion
 CC channel subunits (ICS) has undergone a mutation event so that a cDNA
 CC derived from the subject has any of 134 nucleotide sequences. The method
 CC of the invention has nootropic, neuroprotective, inotropic, antipyretic,
 CC antiarrhythmic, antimigraine, antidepressant, antiparkinsonian,
 CC neuroleptic, tranquilizer, analgesic, nephroprotective, antidiabetic, and
 CC ophthalmological activity. A polynucleotide of the invention acts as an
 CC ion channel agonist, or ion channel antagonist. The methods, isolated
 CC nucleic acids, polypeptides, antibody, selective agonist, antagonist or
 CC modulator of an ion channel, cells and genetically modified non-human
 CC animal, are useful for the diagnosis and treatment of epilepsy and/or a
 CC disorder associated with ion channel dysfunction, such as hyper- or hypo-
 CC kalemic periodic paralysis, myotonia, malignant hyperthermia,
 CC myasthenia, cardiac arrhythmias, episodic ataxia, migraine, Alzheimer's
 CC disease, Parkinson's disease, schizophrenia, hyperkplexia, anxiety,
 CC depression, phobic obsessive symptoms, neuropathic pain, inflammatory
 CC pain, chronic/acute pain, Bartter's syndrome, polycystic kidney disease,
 CC Dent's disease, hyperinsulinaemic hypoglycaemia of infancy, cystic
 CC fibrosis, congenital stationary night blindness and total colour
 CC blindness. The present sequence represents a mutant protein of the
 CC invention. The sequence data for this patent is not represented in the
 CC printed specification, but was obtained in electronic format directly
 CC from WIPO at ftp.wipo.int/pub/published_pat_sequences.
 CC
 XX
 SQ Sequence 2009 AA;
 Query Match 100.0%; Score 28; DB 7; Length 2009;
 Best Local Similarity 100.0%; Pred. No. 7.5e-18;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CLTVFMVMVYIGNLVNLFALLSSSF 28
 Db 968 CLTVFMVMVYIGNLVNLFALLSSSF 995
 RESULT 24
 ADB78594 standard; protein; 2009 AA.
 ID ADB78594
 AC ADB78594;
 XX
 DT 04-DEC-2003 (first entry)
 XX
 DE Human sodium channel subunit mutant SEQ ID NO:138.
 XX
 KW mutein; mutant; ion channel; ion channel subunit; ICS; nootropic;
 KW neuroprotective; inotropic; antipyretic; antiarrhythmic; antimigraine;
 KW antidepressant; antiparkinsonian; neuroleptic; tranquilizer; analgesic;
 KW nephroprotective; antidiabetic; ophthalmological; epilepsy;
 KW ion channel dysfunction; human.
 XX
 OS Synthetic.
 OS Homo sapiens.
 XX
 PN W02003008574-A1.
 XX
 PD 30-JAN-2003.
 XX
 PF 08-JUL-2002; 2002WO-AU000910.

XX
 PR 18-JUL-2001; 2001AU-00006452.
 PR 05-MAR-2002; 2002AU-00000910.
 PR 13-MAY-2002; 2002AU-00002292.
 XX
 PA (BION-) BIONOMICS LTD.
 PA (WALL) WALLACE R W.
 XX
 PI Mulley JC, Harkin LA, Dibbens LM, Phillips HA, Heron SE;
 PI Berkovic SF, Scheffer IE;
 XX
 DR WPI: 2003-239332/23.
 DR N-PSDB; ADB78633.
 PT Identifying predisposition to an ion channel dysfunction, such as
 PT periodic paralysis, cardiac arrhythmias, migraine, Alzheimer's disease,
 PT schizophrenia, anxiety and depression, by detecting encoding-gene
 PT mutation events.
 XX
 PS Claim 13; SEQ ID NO 138; 106pp; English.
 XX
 CC The invention relates to a novel method for identifying a subject
 CC predisposed to a disorder associated with ion channel dysfunction. The
 CC method comprises ascertaining if at least one of the genes encoding ion
 CC channel subunits (ICS) has undergone a mutation event so that a cDNA
 CC derived from the subject has any of 134 nucleotide sequences. The method
 CC of the invention has nootropic, neuroprotective, inotropic, antipyretic,
 CC antiarrhythmic, antimigraine, antidepressant, antiparkinsonian,
 CC neuroleptic, tranquilizer, analgesic, nephroprotective, antidiabetic, and
 CC ophthalmological activity. A polynucleotide of the invention acts as an
 CC ion channel agonist, or ion channel antagonist. The methods, isolated
 CC nucleic acids, polypeptides, antibody, selective agonist, antagonist or
 CC modulator of an ion channel, cells and genetically modified non-human
 CC animal, are useful for the diagnosis and treatment of epilepsy and/or a
 CC disorder associated with ion channel dysfunction, such as hyper- or hypo-
 CC kalemic periodic paralysis, myotonia, malignant hyperthermia,
 CC myasthenia, cardiac arrhythmias, episodic ataxia, migraine, Alzheimer's
 CC disease, Parkinson's disease, schizophrenia, hyperkplexia, anxiety,
 CC depression, phobic obsessive symptoms, neuropathic pain, inflammatory
 CC pain, chronic/acute pain, Bartter's syndrome, polycystic kidney disease,
 CC Dent's disease, hyperinsulinaemic hypoglycaemia of infancy, cystic
 CC fibrosis, congenital stationary night blindness and total colour
 CC blindness. The present sequence represents a mutant protein of the
 CC invention. The sequence data for this patent is not represented in the
 CC printed specification, but was obtained in electronic format directly
 CC from WIPO at ftp.wipo.int/pub/published_pat_sequences.
 CC
 XX
 SQ Sequence 2009 AA;
 Query Match 100.0%; Score 28; DB 7; Length 2009;
 Best Local Similarity 100.0%; Pred. No. 7.5e-18;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CLTVFMVMVYIGNLVNLFALLSSSF 28
 Db 968 CLTVFMVMVYIGNLVNLFALLSSSF 995
 RESULT 25
 ADB78598 standard; protein; 2009 AA.
 ID ADB78598
 AC ADB78598;
 XX
 DT 04-DEC-2003 (first entry)
 XX
 DE Human sodium channel subunit mutant SEQ ID NO:142.
 XX
 KW mutein; mutant; ion channel; ion channel subunit; ICS; nootropic;
 KW neuroprotective; inotropic; antipyretic; antiarrhythmic; antimigraine;
 KW antidepressant; antiparkinsonian; neuroleptic; tranquilizer; analgesic;
 KW nephroprotective; antidiabetic; ophthalmological; epilepsy;
 KW ion channel dysfunction; human.

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 27, 2005, 17:36:05 ; Search time 17 Seconds
(without alignments)
158.475 Million cell updates/sec

Title: US-10-608-584-13

Perfect score: 28

Sequence: 1 CLTVFMVMVIGNLVNLFLALLLSF 28

Scoring table: OLIGO

Gapop 60.0 , Gapext 60.0

Searched: 283416 seqs, 96216763 residues

Word size: 0

Total number of hits satisfying chosen parameters: 283416

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Listing first 100 summaries

Database :
1: pir_79:*
2: pir1:*
3: pir3:*
4: pir4:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	28	100.0	200	2	148108 sodium channel alp
2	28	100.0	2005	2	A46269 sodium channel alp
3	28	100.0	2005	2	B25019 sodium channel pro
4	25	89.3	1976	2	I56555 sodium channel alp
5	23	82.1	1977	2	S54771 sodium channel alp
6	22	78.6	1835	2	I54323 sodium channel alp
7	22	78.6	1836	2	I64893 sodium channel alp
8	22	78.6	1836	2	U50648 sodium channel alp
9	22	78.6	1836	2	I51964 sodium channel alp
10	22	78.6	1840	1	CHRTM1 sodium channel pro
11	21	75.0	1951	2	S00320 sodium channel pro
12	21	75.0	1983	2	A60054 sodium channel pro
13	21	75.0	2016	2	A38195 sodium channel pro
14	21	75.0	2019	2	A33966 sodium channel pro
15	18	64.3	1784	2	I43167 sodium channel pro
16	17	60.7	109	2	S72488 sodium channel pro
17	17	60.7	1034	2	S60051 sodium channel alp
18	17	60.7	1034	2	S60060 sodium channel alp
19	17	60.7	1689	2	S72467 sodium channel pro
20	17	60.7	1820	2	A33299 sodium channel pro
21	17	60.7	1993	2	T30902 sodium channel pro
22	17	60.7	2049	2	T43161 sodium channel pro
23	17	60.7	2049	2	S72458 sodium channel pro
24	16	57.1	2009	2	A25019 sodium channel pro
25	14	50.0	1695	2	J50084 voltage-gated sodi
26	12	42.9	1820	1	CHBE sodium channel pro
27	11	39.3	1739	2	A48298 sodium channel hom
28	10	35.7	1321	2	A60165 sodium channel pro
29	10	35.7	1810	2	T31092 probable voltage-g

30	9	32.1	409	2	E89834 hypothetical prote
31	8	32.1	1957	2	S68453 sodium channel pro
32	8	28.6	501	2	I61512 TNF receptor assoc
33	8	28.6	616	2	AR0263 proteinase IV (EC
34	8	28.6	1699	2	T31340 voltage-gated sodi
35	7	25.0	33	2	A36154 benzphetamine N-de
36	7	25.0	68	2	S49412 fibrinogen-binding
37	7	25.0	100	2	S66718 probable membr
38	7	25.0	101	2	S60632 NADH2 dehydrogen
39	7	25.0	104	1	PFH04A platelet factor 4
40	7	25.0	116	2	B89887 hypothetical prote
41	7	25.0	118	2	S42398 hypothetical prote
42	7	25.0	155	2	S60647 NADH2 dehydrogen
43	7	25.0	166	2	JC6559 interferon-gamma p
44	7	25.0	183	2	E72459 hypothetical prote
45	7	25.0	218	2	T19256 hypothetical prote
46	7	25.0	237	2	AF0772 probable exported
47	7	25.0	247	2	B86161 F1003.11 protein -
48	7	25.0	257	2	H84597 hypothetical prote
49	7	25.0	292	2	A47125 transcription acti
50	7	25.0	358	2	T26231 hypothetical prote
51	7	25.0	386	2	AH3467 glycine betaine/1-
52	7	25.0	390	2	H70904 probable lpxr prot
53	7	25.0	420	2	T39523 probable transmemb
54	7	25.0	451	2	H89798 conserved hypothet
55	7	25.0	480	2	G70302 internalin like pr
56	7	25.0	505	2	AC1469 conserved hypothet
57	7	25.0	513	2	A71004 hypothetical prote
58	7	25.0	538	2	G72539 probable CTP synth
59	7	25.0	578	2	AH1020 cyclochrome c-type
60	7	25.0	721	2	F87611 TonB-dependent rec
61	7	25.0	1158	2	S57348 nuclear factor RIP
62	7	25.0	1172	2	T36053 probable ABC-type
63	7	25.0	1438	2	B71610 Wd40 WEB-1 homolog
64	7	25.0	1582	2	A56248 sulfonylurea recep
65	7	25.0	1765	2	T42368 sodium channel alp
66	6	21.4	52	2	B96798 hypothetical prote
67	6	21.4	66	2	C97928 hypothetical prote
68	6	21.4	73	2	B83338 hypothetical prote
69	6	21.4	83	2	T03673 pili protein (clon
70	6	21.4	85	2	H83492 hypothetical prote
71	6	21.4	97	2	B95038 preprotein translo
72	6	21.4	97	2	H97908 conserved hypothet
73	6	21.4	100	1	F70309 protein export mem
74	6	21.4	102	2	C63655 probable 10kd chap
75	6	21.4	102	2	A75417 hypothetical prote
76	6	21.4	103	2	C71189 hypothetical prote
77	6	21.4	107	2	F71700 hypothetical prote
78	6	21.4	111	2	B70035 chaperonin homolog
79	6	21.4	117	2	AG3451 murein hydrolase e
80	6	21.4	120	2	S57057 probable membrane
81	6	21.4	123	2	A75273 hypothetical prote
82	6	21.4	123	2	T35812 probable small hyd
83	6	21.4	138	2	AD1850 hypothetical prote
84	6	21.4	139	2	B69744 hypothetical prote
85	6	21.4	149	2	F75327 hypothetical prote
86	6	21.4	158	2	G82494 conserved hypothet
87	6	21.4	164	2	S22204 photosystem I chai
88	6	21.4	165	2	B75502 hypothetical prote
89	6	21.4	166	1	IYBOG interferon gamma p
90	6	21.4	166	2	S12723 interferon gamma p
91	6	21.4	166	2	S35795 androgen receptor
92	6	21.4	167	2	D75636 hypothetical prote
93	6	21.4	175	2	B95412 hypothetical prote
94	6	21.4	178	2	C72025 conserved hypothet
95	6	21.4	178	2	A86598 C718 hypothetical
96	6	21.4	180	2	T15426 hypothetical prote
97	6	21.4	184	2	T38162 hypothetical RBR1-
98	6	21.4	184	2	AB2503 hypothetical prote
99	6	21.4	185	2	T24345 hypothetical prote
100	6	21.4	185	2	S76706 hypothetical prote

ALIGNMENTS

RESULT 1

I48108

sodium channel alpha subunit - long-tailed hamster (fragment)

C/Species: Cricetus longicaudatus (long-tailed hamster)

C/Date: 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change 09-Jul-2004

C/Accession: I48108

R/Lalik, P.H.; Krafte, D.S.; Ciccarelli, R.B.

Am. J. Physiol. 264, 803-809, 1993

A/Title: Characterization of endogenous Sodium channel gene expressed in chinese hamster

A/Reference number: I48107

A/Accession: I48108

A/Status: preliminary; translated from GB/EMBL/DBJ

A/Molecule type: mRNA

A/Residues: 1-200 <RES>

A/Cross-references: UNIPROT:Q60464; GB:M87541; NID:g191069; PIDN:AAA36979.1; PID:G553840

C/Genetics:

A/Gene: CNOL

C/Superfamily: sodium channel protein

C/Keywords: duplication

Query Match

Best Local Similarity 100.0%; Score 28; DB 2; Length 200;

Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 45 CLTVFMVWVIGNLVNLFLALLSSSF 72

RESULT 2

A46269

sodium channel alpha chain HBA - human

C/Species: Homo sapiens (man)

C/Date: 20-Oct-1993 #sequence_revision 18-Nov-1994 #text_change 21-Nov-1997

C/Accession: A46269

R/Alamed, C.M.; Ware, D.H.; Lee, S.C.; Patten, C.D.; Ferrer-Montiel, A.V.; Schinder, A.F.

Proc. Natl. Acad. Sci. U.S.A. 89, 8220-8224, 1992

A/Title: Primary structure, chromosomal localization, and functional expression of a vol

A/Reference number: A46269; MUID:92290418; PMID:1325650

A/Accession: A46269

A/Molecule type: mRNA

A/Residues: 1-2005 <AAH>

A/Cross-references: GB:M94055

A/Experimental source: brain

A/Note: sequence extracted from NCBI backbone (NCBIP:113082)

C/Genetics:

A/Map position: 2623-q24.3

C/Superfamily: sodium channel protein

C/Keywords: duplication

Query Match 100.0%; Score 28; DB 2; Length 2005;

Best Local Similarity 100.0%; Pred. No. 2,4e-18; Mismatches 0; Indels 0; Gaps 0;

Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 CLTVFMVWVIGNLVNLFLALLSSSF 28

959 CLTVFMVWVIGNLVNLFLALLSSSF 986

RESULT 3

B25019

sodium channel protein II - rat

C/Species: Rattus norvegicus (Norway rat)

C/Date: 30-Jun-1988 #sequence_revision 30-Jun-1988 #text_change 09-Jul-2004

C/Accession: B25019; S24804

R/Noda, M.; Ikeda, T.; Kayano, T.; Suzuki, H.; Takehana, H.; Kurasaki, M.; Takahashi, H.

Nature 320, 188-192, 1986

A/Title: Existence of distinct sodium channel messenger RNAs in rat brain.

A/Reference number: A93377; MUID:86146901; PMID:3754035

A/Accession: B25019

A/Molecule type: mRNA

A/Residues: 1-2005 <NOD>

A/Cross-references: UNIPROT:Q63509

A/Experimental source: brain

R/Satoh, R.; Gupta, S.K.; Auld, V.J.; Dunn, R.J.

Submitted to the EMBL Data Library, August 1991

A/Description: Developmentally regulated RNA splicing of rat brain sodium channel mRNAs.

A/Reference number: S24803

A/Accession: S24804

A/Status: preliminary

A/Molecule type: DNA

A/Residues: 183-188, 'D', 190-305 <SAR>

A/Cross-references: EMBL:X61149; NID:957074; PIDN:CAA43458.1; PID:G57076

C/Superfamily: sodium channel protein

C/Keywords: duplication; ion transport; sodium channel; transmembrane protein; voltage-g

Query Match

Best Local Similarity 100.0%; Score 28; DB 2; Length 2005;

Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 CLTVFMVWVIGNLVNLFLALLSSSF 28

959 CLTVFMVWVIGNLVNLFLALLSSSF 986

RESULT 4

I56555

sodium channel protein 6 - rat

C/Species: Rattus norvegicus (Norway rat)

C/Date: 26-Jul-1996 #sequence_revision 26-Jul-1996 #text_change 09-Jul-2004

C/Accession: I56555

R/Schaller, K.L.; Krzemien, D.M.; Yarowsky, P.J.; Krueger, B.K.; Caldwell, J.H.

J. Neurosci. 15, 3231-3242, 1995

A/Title: A novel, abundant sodium channel expressed in neurons and glia.

A/Reference number: I56555; MUID:95271284; PMID:7751906

A/Accession: I56555

A/Status: preliminary; translated from GB/EMBL/DBJ

A/Molecule type: mRNA

A/Residues: 1-1976 <RES>

A/Cross-references: UNIPROT:Q63541; GB:U39018; NID:9829033; PIDN:AA642059.1; PID:9829034

C/Genetics:

A/Gene: SC26

C/Superfamily: sodium channel protein

C/Keywords: duplication

Query Match

Best Local Similarity 100.0%; Score 25; DB 2; Length 1976;

Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 4 VFMVWVIGNLVNLFLALLSSSF 28

952 VFMVWVIGNLVNLFLALLSSSF 976

RESULT 5

S54771

sodium channel alpha subunit - human

C/Species: Homo sapiens (man)

C/Date: 27-Oct-1995 #sequence_revision 03-Nov-1995 #text_change 09-Jul-2004

C/Accession: S54771

R/Krugbauer, N.; Lacinova, L.; Flocke, V.; Hofmann, F.

EMBO J. 14, 1084-1090, 1995

A/Title: Structure and functional expression of a new member of the tetrodotoxin-sensitive

A/Reference number: S54771; MUID:95237189; PMID:7720699

A/Accession: S54771

A/Status: preliminary; nucleic acid sequence not shown

A/Molecule type: mRNA

A/Residues: 1-1977 <KLU>

A/Cross-references: UNIPROT:Q15858; EMBL:X82835; NID:g758109; PIDN:CAA58042.1; PID:g7581

C/Superfamily: sodium channel protein

C/Keywords: duplication

Query Match

Best Local Similarity 82.1%; Score 23; DB 2; Length 1977;

Best Local Similarity 100.0%; Pred. No. 1,2e-13;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 6 MMWVIGNLVVLNLFLLALLSSP 28
Db 938 MMWVIGNLVVLNLFLLALLSSP 960

RESULT 6
154323
sodium channel alpha subunit - human
C:Species: Homo sapiens (man)
C:Date: 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change 20-Aug-1999
C:Accession: 154323
R:McClatchey, A.I.; Lin, C.S.; Wang, J.; Hoffman, E.P.; Rojas, C.; Gussella, J.F.
Hum. Mol. Genet. 1, 521-527, 1992
A:Title: The genomic structure of the human skeletal muscle sodium channel gene.
A:Reference number: 154323; MUID:9338444; PMID:1339144
A:Accession: 154323
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-1835 <RES>
A:Cross-references: GB:L01983; NID:G337992; PIDN:AAA75557.1; PID:G908809
C:Genetics:
A:Gene: GDB:SCN4A
A:Cross-references: GDB:125181; OMIM:170500
A:Map position: 17q23.1-17q25.3
A:Interons: 91/3; 131/2; 161/2; 204/2; 235/1; 346/1; 367/2; 414/3; 484/3; 536/1; 615/3; 6
C:Superfamily: sodium channel protein
C:Keywords: duplication

Query March 78.6%; Score 22; DB 2; Length 1835;
Best Local Similarity 100.0%; Pred. No. 9,7e-13;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 7 MMWVIGNLVVLNLFLLALLSSP 28
Db 784 MMWVIGNLVVLNLFLLALLSSP 805

RESULT 7
164893
sodium channel alpha subunit - human
C:Species: Homo sapiens (man)
C:Date: 06-Sep-1996 #sequence_revision 06-Sep-1996 #text_change 09-Jul-2004
C:Accession: 164893
R:George, A.L.
Ann. Neurol. 31, 131-137, 1992
A:Title: Primary structure of the adult human skeletal muscle voltage-dependent sodium c
A:Reference number: 151964; MUID:92246457; PMID:1315496
A:Accession: 164893
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-1836 <RES>
A:Cross-references: UNIPROT:P35499; GB:M81758; NID:G338212; PIDN:AAA60554.1; PID:G338213
C:Genetics:
A:Gene: SKM1
C:Superfamily: sodium channel protein
C:Keywords: duplication

Query March 78.6%; Score 22; DB 2; Length 1836;
Best Local Similarity 100.0%; Pred. No. 9,7e-13;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 7 MMWVIGNLVVLNLFLLALLSSP 28
Db 784 MMWVIGNLVVLNLFLLALLSSP 805

RESULT 8
US0648
sodium channel alpha chain - human
C:Species: Homo sapiens (man)

C:Date: 30-Jun-1992 #sequence_revision 30-Jun-1992 #text_change 09-Jul-2004
C:Accession: US0648; A42099
R:Wang, J.; Rojas, C.V.; Zhou, J.; Schwartz, L.S.; Nicholas, H.; Hoffmann, E.P.
Biochem. Biophys. Res. Commun. 182, 794-801, 1992
A:Title: Sequence and genomic structure of the human adult skeletal muscle sodium channe
A:Reference number: US0648; MUID:92134303; PMID:1310396
A:Accession: US0648
A:Status: nucleic acid sequence not shown
A:Molecule type: mRNA
A:Residues: 1-1836 <MAN>

A:Cross-references: UNIPROT:P35499
A:Note: 861-ASP was also found as the result of polymorphism
R:McClatchey, A.I.; Van den Berg, P.; Perlick-Vance, M.A.; Kaskind, W.; Verellen, C.; M
Cell 68, 769-774, 1992
A:Title: Temperature-sensitive mutations in the III-IV cytoplasmic loop region of the sk
A:Reference number: A42099; MUID:92154689; PMID:1310898
A:Accession: A42099
A:Molecule type: DNA
A:Residues: 1299-1351 <MCC>
A:Cross-references: GB:S82622; NID:G245611; PIDN:AA821450.1; PID:G245612
A:Experimental source: skeletal muscle
A:Note: sequence extracted from NCBI backbone (NCBI:82622, NCBI:P:82623)
C:Genetics:
A:Gene: GDB:SCN4A
A:Cross-references: GDB:125181; OMIM:170500
A:Map position: 17q23.1-17q25.3
C:Superfamily: sodium channel protein
C:Keywords: duplication; glycoprotein; phosphoprotein; transmembrane protein

F:1129-150/Domain: transmembrane #status predicted <TR1>
F:1159-178/Domain: transmembrane #status predicted <TR2>
F:1191-210/Domain: transmembrane #status predicted <TR3>
F:1217-236/Domain: transmembrane #status predicted <TR4>
F:1253-266/Domain: transmembrane #status predicted <TR5>
F:1424-449/Domain: transmembrane #status predicted <TR6>
F:1574-597/Domain: transmembrane #status predicted <R11>
F:1609-632/Domain: transmembrane #status predicted <R12>
F:1641-660/Domain: transmembrane #status predicted <R13>
F:1667-686/Domain: transmembrane #status predicted <R14>
F:1702-724/Domain: transmembrane #status predicted <R15>
F:1777-802/Domain: transmembrane #status predicted <R16>
F:1027-1049/Domain: transmembrane #status predicted <R11>
F:1064-1089/Domain: transmembrane #status predicted <R12>
F:1096-1116/Domain: transmembrane #status predicted <R13>
F:1122-1143/Domain: transmembrane #status predicted <R14>
F:1163-1184/Domain: transmembrane #status predicted <R15>
F:1269-1295/Domain: transmembrane #status predicted <R16>
F:1349-1372/Domain: transmembrane #status predicted <R11>
F:1384-1407/Domain: transmembrane #status predicted <R12>
F:1414-1437/Domain: transmembrane #status predicted <R13>
F:1447-1469/Domain: transmembrane #status predicted <R14>
F:1485-1507/Domain: transmembrane #status predicted <R15>
F:1574-1596/Domain: transmembrane #status predicted <R16>
F:121,149,220,378,415,1019,1130,1242,1313,1721,1826/Binding site: phosphate (Thr) (covalen
F:156,251,513,653,1511,1746/Binding site: phosphate (Ser) (covalent) (by protein kinase A
F:246,288,291,297,303,315,362,507,702,961,1191,1205/Binding site: carboxylate (by protei
F:324,670,725,850,950,1127,1193,1328/Binding site: phosphate (Ser) (covalent) (by protei
F:387,457/Binding site: phosphate (Thr) (covalent) (by protein kinase A) #status predict

Query March 78.6%; Score 22; DB 2; Length 1836;
Best Local Similarity 100.0%; Pred. No. 9,7e-13;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 7 MMWVIGNLVVLNLFLLALLSSP 28
Db 784 MMWVIGNLVVLNLFLLALLSSP 805

RESULT 9
151964
sodium channel alpha chain SCN4A, skeletal muscle - human
C:Species: Homo sapiens (man)
C:Date: 24-May-1996 #sequence_revision 24-May-1996 #text_change 09-Jul-2004
C:Accession: 151964

R.George, A.L.
Ann. Neurol. 31, 131-137, 1992
A:Title: Primary structure of the adult human skeletal muscle voltage-dependent sodium C
A:Accession: I51964; MUID:92246457; PMID:1315496
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-1836 <RES>
A:Cross-references: UNIPROT:P35499; GB:L04236; NID:G292485; PIDN:AA59624.1; PID:G292487
A:Gene: GDB:SCN4A
A:Cross-references: GDB:125181; OMIM:170500
A:Map position: 17q23.1-17q25.3
A:Introns: 91/3; 131/2; 161/2; 204/2; 235/1; 346/1; 367/2; 414/3; 484/3; 536/1; 615/3; 6
C:Superfamily: sodium channel protein
C:Keywords: duplication; skeletal muscle

Query Match 78.6%; Score 22; DB 2; Length 1836;
Best Local Similarity 100.0%; Pred. No. 9.7e-13;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 7 MWVIGNLVNLFLALLSSP 28
DB 784 MWVIGNLVNLFLALLSSP 805

RESULT 10
CHRTM1
sodium channel protein mul alpha chain, skeletal muscle - rat
C:Species: Rattus norvegicus (Norway rat)
C:Date: 30-Sep-1990 #sequence_revision 30-Sep-1990 #text_change 09-Jul-2004
C:Accession: JN0007
R:Timer: J.S.; Cooperman, S.S.; Tomiko, S.A.; Zhou, J.; Crean, S.M.; Boyle, M.B.; Kall
Neuron 3, 33-49, 1995
A:Title: Primary structure and functional expression of a mammalian skeletal muscle sodi
A:Reference number: JN0007; MUID:90148778; PMID:2559760
A:Accession: JN0007
A:Molecule type: mRNA
A:Residues: 1-1840 <TRI>
A:Cross-references: UNIPROT:P15390; GB:M26643; NID:G205651; PIDN:AAA1682.1; PID:G205652
C:Comment: Action potentials propagated by skeletal muscle sodium channels are responsib
C:Comment: This heavily glycosylated protein contains four homologous domains, each of w
C:Superfamily: sodium channel protein
C:Keywords: duplication; glycoprotein; ion transport; neuromuscular junction; phosphop
F:120-458,561-813,1013-1305,1335-1611/region: duplication
F:56,251,1321,1504/Binding site: phosphate (Ser) (covalent) [by CAMP-dependent kinase] #
F:214,286,291,297,303,309,315,327,356,502,696,954,1184,1198,1563,1702/Binding site: cati

Query Match 78.6%; Score 22; DB 1; Length 1840;
Best Local Similarity 100.0%; Pred. No. 9.7e-13;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 7 MWVIGNLVNLFLALLSSP 28
DB 778 MWVIGNLVNLFLALLSSP 799

RESULT 11
S00320
sodium channel protein III - rat
C:Species: Rattus norvegicus (Norway rat)
C:Date: 30-Jun-1989 #sequence_revision 30-Jun-1989 #text_change 09-Jul-2004
C:Accession: S00320
R:Kayano, T.; Noda, M.; Flockerzi, V.; Takahashi, H.; Numa, S.
FEBS Lett. 228, 187-194, 1988
A:Title: Primary structure of rat brain sodium channel III deduced from the cDNA sequenc
A:Reference number: S00320; MUID:86137594; PMID:2449363
A:Accession: S00320
A:Molecule type: mRNA
A:Residues: 1-1951 <RAY>
A:Cross-references: UNIPROT:P06104; EMBL:Y00766; NID:957210; PIDN:CAA68735.1; PID:957211
A>Note: 270-Ile, 278-Leu, 355-Thr, 513-Lys, and 1059-Arg were also found

C:Superfamily: sodium channel protein
C:Keywords: duplication; transmembrane protein

Query Match 75.0%; Score 21; DB 2; Length 1951;
Best Local Similarity 100.0%; Pred. No. 8.9e-12;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 8 MWVIGNLVNLFLALLSSP 28
DB 918 MWVIGNLVNLFLALLSSP 938

RESULT 12
A60054
sodium channel protein IIIB, long form - rat
C:Species: Rattus norvegicus (Norway rat)
C:Date: 03-Mar-1993 #sequence_revision 03-Mar-1993 #text_change 09-Jul-2004
C:Accession: A60054; B44824
R:Joho, R.H.; Moorman, J.R.; Vandongen, A.M.J.; Kirsch, G.E.; Silberberg, H.; Schuster, C
Brain Res. Mol. Brain Res. 7, 105-113, 1990
A:Title: Toxin and kinetic profile of rat brain type III sodium channels expressed in Xer
A:Reference number: A60054; MUID:90251117; PMID:2160038
A:Accession: A60054
A:Status: not compared with conceptual translation
A:Molecule type: mRNA
A:Residues: 1-1983 <JOH>
A:Cross-references: UNIPROT:Q64243
R:Schaller, K.L.; Kremien, D.W.; McKenna, N.M.; Caldwell, J.H.
J. Neurosci. 12, 1370-1381, 1992
A:Title: Alternatively spliced sodium channel transcripts in brain and muscle.
A:Reference number: A44824; MUID:92211397; PMID:1313493
A:Accession: B44824
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 611-662 <SCH>
A:Cross-references: GB:S97388; NID:G248225; PIDN:AA21984.1; PID:G248226
A:Experimental source: skeletal muscle
A>Note: Sequence inconsistent with the nucleotide translation
A:Note: Sequence extracted from NCBI backbone (NCBIN:97388, NCBI:P:97391)
C:Superfamily: sodium channel protein
C:Keywords: duplication; glycoprotein; ion transport; sodium channel; transmembrane prote

Query Match 75.0%; Score 21; DB 2; Length 1983;
Best Local Similarity 100.0%; Pred. No. 9e-12;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 8 MWVIGNLVNLFLALLSSP 28
DB 950 MWVIGNLVNLFLALLSSP 970

RESULT 13
A38195
sodium channel protein hHT, cardiac - human
N:Alternate names: tetrodotoxin-insensitive, voltage-dependent sodium channel, TTX-I NaC
C:Species: Homo sapiens (man)
C:Date: 31-Dec-1993 #sequence_revision 31-Dec-1993 #text_change 09-Jul-2004
C:Accession: A38195
R:Gellens, M.E.; George Jr., A.L.; Chen, L.O.; Chahine, M.; Horn, R.; Barchi, R.L.; Kall
Proc. Natl. Acad. Sci. U.S.A. 89, 554-558, 1992
A:Title: Primary structure and functional expression of the human cardiac tetrodotoxin-I
A:Reference number: A38195; MUID:92115699; PMID:11309946
A:Accession: A38195
A:Status: nucleic acid sequence not shown
A:Molecule type: mRNA
A:Residues: 1-2016 <GEL>
A:Cross-references: UNIPROT:Q14524; GB:M77235; NID:G184038; PIDN:AAA5644.1; PID:G184039
A:Experimental source: heart
C:Superfamily: sodium channel protein
C:Keywords: cardiac muscle; duplication; glycoprotein; heart; ion transport; sodium chan
Query Match 75.0%; Score 21; DB 2; Length 2016;
Best Local Similarity 100.0%; Pred. No. 9.2e-12;

Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 8 VMVIGNLVVNLFLALLSSSF 28
 |||||
 Db 922 VMVIGNLVVNLFLALLSSSF 942

RESULT 14

sodium channel protein I, cardiac - rat

N/Alternate names: sodium channel protein (SKM2) alpha chain

C/Species: Rattus norvegicus (Norway rat)

C/Date: 30-Mar-1990 #sequence_revision 30-Mar-1990 #text_change 09-Jul-2004

C/Accession: A33996; J00412

R/Rogart, R.B.; Cribbs, L.L.; Muglia, L.K.; Kephart, D.D.; Kaiser, M.W.

Proc. Natl. Acad. Sci. U.S.A. 86, 8170-8174, 1989

A/Title: Molecular cloning of a putative tetrodotoxin-resistant rat heart Na(+) channel

A/Reference number: A33996; MUID:90046760; PMID:2554302

A/Accession: A33996

A/Status: preliminary

A/Molecule type: mRNA

A/Residues: 1-2019 <RGS>

A/Cross-references: UNIPROT:P15389; GB:M27902; NID:G206857; PIDN:AAA2114.1; PID:G206858

R/Kallen, R.G.; Sheng, Z.H.; Yang, J.; Chen, L.; Rogart, R.B.; Barchi, R.L.

Neuron 4, 233-242, 1990

A/Title: Primary structure and expression of a sodium channel characteristic of denervat

A/Reference number: J00412; MUID:90166613; PMID:2155010

A/Accession: J00412

A/Molecule type: mRNA

A/Residues: 1-479; 481-1712; 'T', 1714-1963; 'R', 1965-2019 <KAL>

A/Experimental source: muscle

C/KeyWords: sodium channel protein

C/KeyWords: cardiac muscle; duplication; heart; sodium channel; transmembrane protein

Query Match 75.0%; Score 21; DB 2; Length 2019;

Best Local Similarity 100.0%; Pred. No. 9.2e-12;

Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 8 VMVIGNLVVNLFLALLSSSF 28
 |||||
 Db 925 VMVIGNLVVNLFLALLSSSF 945

RESULT 15

sodium channel protein - California market squid

C/Species: Loligo opalescens (California market squid)

C/Date: 11-Jan-2000 #sequence_revision 11-Jan-2000 #text_change 09-Jul-2004

C/Accession: T43167

R/Rosenthal, J.J.; Gilly, W.F.

Proc. Natl. Acad. Sci. U.S.A. 90, 10026-10030, 1993

A/Title: Amino acid sequence of a putative sodium channel expressed in the giant axon of

A/Reference number: 222324; MUID:94052096; PMID:8234251

A/Accession: T43167

A/Status: preliminary

A/Molecule type: mRNA

A/Residues: 1-1784 <ROS>

A/Cross-references: UNIPROT:Q25377; EMBL:L19979; NID:G349118; PID:G349119; PIDN:AAA16202

A/Experimental source: stellate ganglia

C/Suprafamily: sodium channel protein

C/KeyWords: ion transport; membrane protein; sodium channel; voltage-gated ion channel

Query Match 64.3%; Score 18; DB 2; Length 1784;

Best Local Similarity 100.0%; Pred. No. 5.5e-09;

Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 11 IGNLVVNLFLALLSSSF 28
 |||||
 Db 799 IGNLVVNLFLALLSSSF 816

RESULT 16

S72488

sodium channel protein para-type alpha chain - house fly (strain SBO) (fragment)

C/Species: Musca domestica (house fly)

A/Variety: strain SBO

C/Date: 24-Oct-1998 #sequence_revision 24-Oct-1998 #text_change 09-Jul-2004

C/Accession: S72488

R/Miyazaki, M.; Ohnaka, K.; Dunlap, D.Y.; Matsumura, F.

Mol. Gen. Genet. 252, 61-68, 1996

A/Title: Cloning and sequencing of the para-type sodium channel gene from susceptible an

A/Reference number: S72487; MUID:96397510; PMID:8804404

A/Accession: S72488

A/Status: not compared with conceptual translation

A/Molecule type: mRNA

A/Residues: 1-109 <MT>

A/Cross-references: UNIPROT:Q25439; UNIPROT:Q25440; UNIPROT:Q254615; UNIPROT:Q254617

A/Experimental source: strain SBO

C/Suprafamily: sodium channel protein

C/KeyWords: sodium channel; transmembrane protein

Query Match 60.7%; Score 17; DB 2; Length 109;

Best Local Similarity 100.0%; Pred. No. 4.3e-09;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 10 VIGNLVVNLFLALLS 26
 |||||
 Db 71 VIGNLVVNLFLALLS 87

RESULT 17

sodium channel alpha chain, exon c-containing splice form - fruit fly (Drosophila virilis)

C/Species: Drosophila virilis

C/Date: 24-Aug-1996 #sequence_revision 08-Nov-1996 #text_change 09-Jul-2004

C/Accession: S60051

R/Thackeray, J.R.; Ganetzky, B.

Genetics 141, 203-214, 1995

A/Title: Conserved alternative splicing patterns and splicing signals in the Drosophila

A/Reference number: S60051; MUID:96042905; PMID:8536968

A/Accession: S60051

A/Status: nucleic acid sequence not shown

A/Molecule type: nucleic acid

A/Residues: 1-1034 <THA>

A/Cross-references: UNIPROT:Q24714; EMBL:U26343

C/Genetics:

A/Gene: FlyBase:FlyBase:FBgn0015214

A/Cross-references: FlyBase:FBgn0015214

C/Suprafamily: sodium channel protein

C/KeyWords: alternative splicing; duplication; transmembrane protein

F/306-329/Region: alternatively spliced segment 1 (exon 4) #status experimental

F/330-350/Region: alternatively spliced segment a #status experimental

F/538-545/Region: alternatively spliced segment b (exon 9) #status experimental

F/874-886/Region: alternatively spliced segment c (exon 12) #status experimental

F/887-896/Region: alternatively spliced segment d (exon 13) #status experimental

F/958-982/Region: alternatively spliced segment h (exon 14) #status experimental

Query Match 60.7%; Score 17; DB 2; Length 1034;

Best Local Similarity 100.0%; Pred. No. 3e-08;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 10 VIGNLVVNLFLALLS 26
 |||||
 Db 800 VIGNLVVNLFLALLS 816

RESULT 18

sodium channel alpha chain, exon d-containing splice form - fruit fly (Drosophila virilis)

C/Species: Drosophila virilis

C/Date: 24-Aug-1996 #sequence_revision 08-Nov-1996 #text_change 09-Jul-2004

C/Accession: S60060

R/Thackeray, J.R.; Ganetzky, B.

Genetics 141, 203-214, 1995

A/Title: Conserved alternative splicing patterns and splicing signals in the Drosophila

A/Reference number: S60051; MUID:96042905; PMID:8536968

A:Accession: S60060
A:Status: nucleic acid sequence not shown
A:Molecule type: nucleic acid
A:Residues: 1-1034 <THA>
A:Cross-references: UNIPROT:Q24714; EMBL:U26343
C:Genetics:
A:Gene: FlyBase:Dmir/para
A:Cross-references: FlyBase:FBgn0015214
C:Superfamily: sodium channel protein
C:Keywords: alternative splicing; duplication; transmembrane protein
F:306-329/Region: alternatively spliced segment i (exon 4) #status experimental
F:330-350/Region: alternatively spliced segment a #status experimental
F:538-545/Region: alternatively spliced segment b (exon 9) #status experimental
F:874-886/Region: alternatively spliced segment e (exon 12) #status experimental
F:887-896/Region: alternatively spliced segment f (exon 13) #status experimental
F:958-982/Region: alternatively spliced segment h (exon 14) #status experimental

Query Match 60.7%; Score 17; DB 2; Length 1034;
Best Local Similarity 100.0%; Pred. No. 3e-08;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 10 VIGNLVNLFALILS 26
Db 800 VIGNLVNLFALILS 816

RESULT 19
S72467
sodium channel protein para-type alpha chain - German cockroach (strain CSMA) (fragment)
C:Species: Blattella germanica (German cockroach)
A:Variety: strain CSMA
C>Date: 29-Jul-1997 #sequence_revision 29-Aug-1997 #text_change 09-Jul-2004
A:Accession: S72467; S72487
R:MiYazaki, M.; Ohyama, K.; Dunlap, D.Y.; Matsumura, F.
submitted to the EMBL Data Library, September 1996
A:Description: Cloning and sequencing of the para-type sodium channel gene from susceptible
A:Reference number: S72467
A:Accession: S72467
A:Molecule type: mRNA
A:Residues: 1-1689 <MTY>
A:Cross-references: UNIPROT:Q93135; EMBL:U71083; NID:G1633647; PIDN:AA82037.1; PID:G163
R:MiYazaki, M.; Ohyama, K.; Dunlap, D.Y.; Matsumura, F.
Mol. Genet. 252, 61-68, 1996
A:Title: Cloning and sequencing of the para-type sodium channel gene from susceptible an
A:Reference number: S72487; MUID:96397510; PMID:8804404
A:Accession: S72487
A:Molecule type: mRNA
A:Residues: 711-819 <MTW>
A:Cross-references: EMBL:U71083
C:Superfamily: sodium channel protein
C:Keywords: duplication; sodium channel; transmembrane protein

Query Match 60.7%; Score 17; DB 2; Length 1689;
Best Local Similarity 100.0%; Pred. No. 4.5e-08;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 10 VIGNLVNLFALILS 26
Db 781 VIGNLVNLFALILS 797

RESULT 20
A33299
sodium channel protein - fruit fly (Drosophila melanogaster) (fragment)
C:Species: Drosophila melanogaster
C>Date: 20-Dec-1989 #sequence_revision 20-Dec-1989 #text_change 21-Nov-1997
A:Accession: A33299
R:Loughney, K.; Kreber, R.; Ganetzky, B.
Cell 58, 1143-1154, 1989
A:Title: Molecular analysis of the para locus, a sodium channel gene in Drosophila.
A:Reference number: A33299; MUID:89376565; PMID:2550145
A:Accession: A33299
A:Status: preliminary

A:Molecule type: mRNA
A:Residues: 1-1820 <LOU>
A:Cross-references: GB:M32078; GB:M24285
C:Genetics:
A:Gene: FlyBase:para
A:Cross-references: FlyBase:FBgn0003036
C:Superfamily: sodium channel protein
C:Keywords: duplication; phosphoprotein; transmembrane protein

Query Match 60.7%; Score 17; DB 2; Length 1820;
Best Local Similarity 100.0%; Pred. No. 4.8e-08;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 10 VIGNLVNLFALILS 26
Db 1004 VIGNLVNLFALILS 1020

RESULT 21
T30902
sodium channel SCAP1 alpha chain - California sea hare
C:Species: Aplysia californica (California sea hare)
C>Date: 22-Oct-1999 #sequence_revision 22-Oct-1999 #text_change 09-Jul-2004
A:Accession: T30902
R:Dyer, J.R.; Johnston, W.J.; Castellucci, V.F.; Dunn, R.J.
DNA Cell Biol. 16, 347-356, 1997
A:Title: Cloning and tissue distribution of the Aplysia Na+ channel alpha-subunit cDNA.
A:Reference number: Z20929; MUID:97238630; PMID:9115644
A:Accession: T30902
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-1993 <DYB>
A:Cross-references: UNIPROT:P90670; EMBL:U66915; NID:G1842248; PID:G1842249; PIDN:AA0474;
C:Superfamily: sodium channel protein

Query Match 60.7%; Score 17; DB 2; Length 1993;
Best Local Similarity 100.0%; Pred. No. 5.2e-08;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 12 GNLVNLFLALISSP 28
Db 971 GNLVNLFLALISSP 987

RESULT 22
T43161
sodium channel protein Tuna1 - sea squirt (Halocynthia roretzi)
C:Species: Halocynthia roretzi
C>Date: 11-Jan-2000 #sequence_revision 11-Jan-2000 #text_change 09-Jul-2004
A:Accession: T43161
R:Okamura, Y.; Ono, F.; Okagaki, R.; Chong, J.; Mandel, G.
Neuron 13, 937-948, 1994
A:Title: Neural expression of a sodium channel gene requires cell-specific interactions.
A:Reference number: Z22220; MUID:95033215; PMID:7946338
A:Accession: T43161
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-2049 <OKA>
A:Cross-references: UNIPROT:Q25150; EMBL:U71311; PIDN:BA04133.1
C:Superfamily: sodium channel protein
C:Keywords: sodium channel; transmembrane protein

Query Match 60.7%; Score 17; DB 2; Length 2049;
Best Local Similarity 100.0%; Pred. No. 5.4e-08;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 12 GNLVNLFLALISSP 28
Db 1009 GNLVNLFLALISSP 1025

RESULT 23
S72458

sodium channel protein para-type alpha chain - house fly (strain Cooper)
C:Species: Musca domestica (house fly)
A:Variety: strain Cooper
C:Date: 24-Oct-1998 #sequence_revision 24-Oct-1998 #text_change 09-Jul-2004
C:Accession: S72458
R:Williamson, M.S.; Martinez-Torres, D.; Hick, C.A.; Devonshire, A.L.
Mol. Gen. Genet. 252, 51-60, 1996
A:Title: Identification of mutations in the housefly para-type sodium channel gene assoc
A:Reference number: S72458; MUID:96397509; PMID:8804403
A:Accession: S72458
A:Molecule type: mRNA
A:Residues: 1-2108 <NID>
A:Cross-references: UNIPROT:Q94615; EMBL:X96668
A:Experimental source: strain Cooper
C:Genetics:
A:Map position: 3
C:Superfamily: sodium channel protein
C:Keywords: alternative splicing; glycoprotein; phosphoprotein; sodium channel; transmem
F:302,314,332,367,1451,1470/Binding site: carbohydrate (Asn) (covalent) #status predicted
F:541,1208,1582/Binding site: phosphate (Ser) (covalent) #status predicted

Query Match 60.7%; Score 17; DB 2; Length 2108;
Best Local Similarity 100.0%; Pred. No. 5.5e-08;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 10 VIGNLVNLFLLALLS 26
DB 1010 VIGNLVNLFLLALLS 1026

RESULT 24
A25019

sodium channel protein I - rat
N:Alternate names: sodium channel protein A

C:Species: Rattus norvegicus (Norway rat)

C:Date: 30-Jun-1988 #sequence_revision 30-Jun-1988 #text_change 09-Jul-2004

C:Accession: A25019; S40783; I84764

R:Noda, M.; Ikeda, T.; Kayano, T.; Suzuki, H.; Takeshima, H.; Kurasaki, M.; Takahashi, H

Nature 320, 188-192, 1986

A:Title: Existence of distinct sodium channel messenger RNAs in rat brain.

A:Reference number: A33377; MUID:86146901; PMID:3754035

A:Accession: A25019

A:Molecule type: mRNA

A:Residues: 1-2009 <NOD>

A:Cross-references: UNIPROT:P04774; GB:X03638; NID:G57216; PIDN:CAA27286.1; PID:G57217

A:Experimental source: brain

R:Sarao, R.; Gupta, S.K.; Auld, V.J.; Dunn, R.J.

Nucleic Acids Res. 19, 5673-5679, 1991

A:Title: Developmentally regulated alternative RNA splicing of rat brain sodium channel

A:Reference number: S40782; MUID:9205114; PMID:1658739

A:Accession: S40783

A:Molecule type: DNA

A:Residues: 177-253 <SAR>

R:Noda, M.; Numa, S.

J. Recept. Res. 7, 467-497, 1987

A:Title: Structure and function of sodium channel.

A:Reference number: I50536; MUID:87311395; PMID:2442385

A:Accession: I84764

A:Status: preliminary; translated from GB/EMBL/DBJ

A:Molecule type: mRNA

A:Residues: 1-2009 <RES>

A:Cross-references: GB:M22253; NID:G1041088; PIDN:AAA79965.1; PID:G1041089

C:Superfamily: sodium channel protein

C:Keywords: duplication; ion transport; sodium channel; transmembrane protein; voltage-g

Query Match 57.1%; Score 16; DB 2; Length 2009;
Best Local Similarity 100.0%; Pred. No. 4.6e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 13 NLVNTLFLALLSSF 28
DB 980 NLVNTLFLALLSSF 995

RESULT 25
J50084
voltage-gated sodium channel alpha subunit - hydromedusa (Polyorchis penicillatus)

N:Alternate names: PpSCN 1

C:Species: Polyorchis penicillatus

C:Date: 11-May-1998 #sequence_revision 29-May-1998 #text_change 09-Jul-2004

C:Accession: J50084

R:Spafford, J.D.; Spencer, A.N.; Gallin, W.J.

Biochem. Biophys. Res. Commun. 244, 772-780, 1998

A:Title: A putative voltage-gated sodium channel alpha subunit (PpSCN1) from the hydromedusa

A:Reference number: J50084; MUID:98205197; PMID:9535741

A:Accession: J50084

A:Molecule type: mRNA

A:Residues: 1-1695 <SPA>

A:Cross-references: UNIPROT:O62604; GB:AF047380; NID:G3005563; PIDN:AAC38974.1; PID:G300

C:Comment: This protein is the only pore-forming alpha subunit available to account for

C:Superfamily: sodium channel protein

C:Keywords: glycoprotein

F:201,273,299,684,1065,1082,1089,1428/Binding site: carbohydrate (Asn) (covalent) #status

Query Match 50.0%; Score 14; DB 2; Length 1695;
Best Local Similarity 100.0%; Pred. No. 3e-05;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 15 VVNLFLALLSSF 28
DB 704 VVNLFLALLSSF 717

Search completed: January 27, 2005, 17:52:44
Job time : 19 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 CompuGen Ltd.

OM protein - protein search, using SW model

Run on: January 27, 2005, 17:35:20 ; Search time 92.5 Seconds
(without alignments)
174.167 Million cell updates/sec

Title: US-10-608-584-13

Perfect score: 28
Sequence: 1 CLTFFMMVIGNLVNLFLALLLSF 28

Scoring table: OLIGO
Gapop 60.0 , Gapext 60.0

Searched: 1825181 seqs, 575374646 residues

Word size : 0

Total number of hits satisfying chosen parameters: 1825181

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Listing first 100 summaries

Database : UniProt_02:*
1: uniprot_sprot:*
2: uniprot_trembl:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match Length	ID	Description
1	28	100.0	200 2 Q60464	Q60464 cricetus
2	28	100.0	276 2 Q8UVC2	Q8UVC2 pagotus
3	28	100.0	500 2 Q800U0	Q800U0 ictalurus p
4	28	100.0	1981 2 Q8UUV6	Q8UUV6 homo sapien
5	28	100.0	2005 1 C1N2_HUMAN	Q92550 homo sapien
6	28	100.0	2005 1 C1N2_RAT	P04775 cynops pyr
7	28	100.0	2007 2 Q9YGN7	Q9YGN7 cynops pyr
8	28	100.0	2009 1 C1N1_HUMAN	P35498 homo sapien
9	25	89.3	472 2 Q800U4	Q800U4 ictalurus p
10	25	89.3	743 2 Q90Z26	Q90Z26 sternopygus
11	25	89.3	751 2 Q90Z28	Q90Z28 sternopygus
12	25	89.3	1717 2 Q90519	Q90519 fuqu rubrip
13	25	89.3	1880 2 Q91BF1	Q91BF1 takifugu pa
14	25	89.3	1949 2 Q9DF53	Q9DF53 brachydact
15	25	89.3	1976 2 Q63541	Q63541 ratius norv
16	25	89.3	1978 1 C1N8_MOUSE	Q9WCU3 mus musculu
17	25	89.3	1978 1 Q88420	Q88420 ratius norv
18	25	89.3	1980 1 C1N8_HUMAN	Q9UQD0 homo sapien
19	25	89.3	1988 1 Q88421	Q88421 ratius norv
20	23	82.1	1977 2 Q15858	Q15858 homo sapien
21	23	82.1	1984 2 Q28644	Q28644 oryctolagus
22	23	82.1	1984 2 Q08562	Q08562 ratius norv
23	22	78.6	473 2 Q800T9	Q800T9 ictalurus p
24	22	78.6	488 2 Q800U1	Q800U1 ictalurus p
25	22	78.6	744 2 Q90Z27	Q90Z27 sternopygus
26	22	78.6	1836 1 C1N4_HUMAN	P35499 homo sapien
27	22	78.6	1840 1 C1N4_RAT	P15390 ratius norv
28	22	78.6	1841 2 Q70611	Q70611 ratius norv
29	21	75.0	289 2 Q9ER60	Q9ER60 mus musculu
30	21	75.0	289 2 Q62204	Q62204 mus musculu
31	21	75.0	479 2 Q800U3	Q800U3 ictalurus p

32	21	75.0	488 2	Q800T8	Q800T8 ictalurus p
33	21	75.0	588 2	Q8U667	Q8U667 xenopus lae
34	21	75.0	1136 2	Q804F4	Q804F4 sternopygus
35	21	75.0	1951 1	C1N3_RAT	P08104 ratius norv
36	21	75.0	1951 1	Q9C007	Q9C007 homo sapien
37	21	75.0	1962 2	Q7SRX3	Q7SRX3 homo sapien
38	21	75.0	1962 2	BAD12085	BAD12085 homo sapi
39	21	75.0	1966 2	Q925G6	Q925G6 ratius norv
40	21	75.0	2000 1	C1N3_HUMAN	Q9Y456 homo sapien
41	21	75.0	2013 2	Q6E5M3	Q6E5M3 canis faml
42	21	75.0	2015 2	Q6ETR3	Q6ETR3 homo sapien
43	21	75.0	2015 2	Q81ZC9	Q81ZC9 homo sapien
44	21	75.0	2015 2	Q96J69	Q96J69 homo sapien
45	21	75.0	2016 1	C1N5_HUMAN	Q14524 homo sapien
46	21	75.0	2016 2	Q75RY0	Q75RY0 homo sapien
47	21	75.0	2016 2	BAD12084	BAD12084 homo sapi
48	21	75.0	2019 1	C1N5_RAT	P15389 ratius norv
49	21	75.0	2019 1	Q9UJY9	Q9UJY9 mus musculu
50	21	75.0	2022 2	Q8WMP8	Q8WMP8 bos tauru
51	20	71.4	742 2	Q90Z29	Q90Z29 sternopygus
52	20	67.9	1089 2	Q81S97	Q81S97 varroa deat
53	19	67.9	1130 2	Q9XZC1	Q9XZC1 boophilus m
54	19	67.9	2215 2	Q8ED77	Q8ED77 varroa deat
55	18	64.3	1784 2	Q25377	Q25377 loligo opal
56	17	60.7	47 2	Q6TFE2	Q6TFE2 anopheles s
57	17	60.7	47 2	Q6V9Y7	Q6V9Y7 anopheles f
58	17	60.7	47 2	Q6VE68	Q6VE68 anopheles c
59	17	60.7	47 2	Q6VE70	Q6VE70 anopheles c
60	17	60.7	47 2	Q7JNL8	Q7JNL8 heliochis v
61	17	60.7	47 2	AAK32092	AAK32092 anopheles
62	17	60.7	47 2	AAK32093	AAK32093 anopheles
63	17	60.7	47 2	AAK32094	AAK32094 anopheles
64	17	60.7	47 2	AAK32095	AAK32095 anopheles
65	17	60.7	47 2	AAK32095	AAK32095 anopheles
66	17	60.7	47 2	AAK32095	AAK32095 anopheles
67	17	60.7	124 2	Q86LC6	Q86LC6 melligethes
68	17	60.7	130 2	Q8MU84	Q8MU84 aedes aegy
69	17	60.7	132 2	Q8IT45	Q8IT45 pediculus h
70	17	60.7	132 2	Q8IT46	Q8IT46 pediculus h
71	17	60.7	136 2	Q45207	Q45207 anopheles g
72	17	60.7	140 2	Q6KBA2	Q6KBA2 ctenocephal
73	17	60.7	140 2	Q95VK8	Q95VK8 aphid gossy
74	17	60.7	140 2	Q95VK9	Q95VK9 aphid gossy
75	17	60.7	140 2	Q95VK9	Q95VK9 aphid gossy
76	17	60.7	142 2	Q97167	Q97167 culex pipie
77	17	60.7	162 2	Q710W0	Q710W0 franklini
78	17	60.7	162 2	Q710W1	Q710W1 franklini
79	17	60.7	162 2	CAC79234	CAC79234 franklini
80	17	60.7	162 2	CAC79235	CAC79235 franklini
81	17	60.7	162 2	CAC79236	CAC79236 franklini
82	17	60.7	220 2	Q95VPA	Q95VPA aphid gossy
83	17	60.7	316 2	Q18460	Q18460 haematobia
84	17	60.7	329 2	Q18461	Q18461 haematobia
85	17	60.7	335 2	Q7JP67	Q7JP67 drosophila
86	17	60.7	340 2	Q7JP66	Q7JP66 drosophila
87	17	60.7	348 2	Q24726	Q24726 drosophila
88	17	60.7	508 2	Q800U2	Q800U2 ictalurus p
89	17	60.7	576 2	Q6DLU1	Q6DLU1 aedes aegy
90	17	60.7	603 2	Q6DLT6	Q6DLT6 aedes albop
91	17	60.7	626 2	Q6DLT5	Q6DLT5 aedes albop
92	17	60.7	660 2	Q24717	Q24717 drosophila
93	17	60.7	689 2	Q24715	Q24715 drosophila
94	17	60.7	693 2	Q24712	Q24712 drosophila
95	17	60.7	701 2	Q24713	Q24713 drosophila
96	17	60.7	704 2	Q24716	Q24716 drosophila
97	17	60.7	706 2	Q24711	Q24711 drosophila
98	17	60.7	714 2	Q24714	Q24714 drosophila
99	17	60.7	1087 2	Q9XYM6	Q9XYM6 leptinotars
100	17	60.7	1347 2	Q7PMT4	Q7PMT4 anopheles g

ALIGNMENTS

```

RESULT 1
Q0464 ID 060464 PRELIMINARY; PRT; 200 AA.
AC 060464;
DT 01-NOV-1996 (TREMBlrel. 01, Created)
DT 01-NOV-1996 (TREMBlrel. 01, last sequence update)
DE 01-MAR-2004 (TREMBlrel. 26, last annotation update)
DE Sodium channel alpha subunit (Fragment).
GN Name=choi;
OS Cricetus longicaudatus (Long-tailed hamster) (Chinese hamster).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Cricetinae;
OC Cricetus.
OC NCBI_TaxID=10030;
RN [1]
RP SEQUENCE FROM N.A.
RX TISSUE=Ovary;
MEDLINE=93235989; PubMed=7682773;
RA Lalik P.H., Krafte D.S., Ciccarelli R.B.;
RT "Characterization of endogenous Sodium channel gene expressed in
RL Chinese hamster ovary cells";
RL Am. J. Physiol. 264:R03-R09(1993).
DR EMBL: M87541; AAA36979.1; -.
DR PIR: I48108; I48108.
DR GO: GO:0016021; C:Integral to membrane; IEA.
DR GO: GO:0005216; F:Ion channel activity; IEA.
DR GO: GO:0006811; P:Ion transport; IEA.
DR InterPro: IPR005821; Ion trans.
DR InterPro: IPR010526; Na_trans_assoc.
DR Pfam: PF06512; Ion_trans_1.
DR Pfam: PF06512; Na_trans_assoc; 1.
DR Ion transport; Ionic channel; Transmembrane; Transport.
KW NON_TER 200
FT SEQUENCE 200 AA; 22676 MW; A09791C08E43458 CRC64;
SQ

Query Match 100.0%; Score 28; DB 2; Length 200;
Best Local Similarity 100.0%; Pred. No. 3.9e-18;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CLTVFMVWVIGNLVNLFLALLSSSF 28
Db 45 CLTVFMVWVIGNLVNLFLALLSSSF 72

RESULT 2
Q8UVC2 PRELIMINARY; PRT; 276 AA.
ID Q8UVC2;
AC Q8UVC2;
DT 01-MAR-2002 (TREMBlrel. 20, Created)
DT 01-MAR-2002 (TREMBlrel. 20, last sequence update)
DE 01-MAR-2004 (TREMBlrel. 26, last annotation update)
DE Putative voltage-activated sodium channel alpha subunit
DE (Fragment).
GN Name=scr4a;
OS Pagothia bernacchi (Emerald rockcod) (Trematomus bernacchi).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neocleostei;
OC Acanthomorpha; Acanthopterygii; Percomorpha; Perciformes;
OC Notocheniidae; Notocheniidae; Trematomus.
OX NCBI_TaxID=40690;
RN [1]
RP SEQUENCE FROM N.A.
RX TISSUE=Skeletal muscle;
Moran O., Elia L.;
RA Submitted (JAN-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF465250; AAL69973.1; -.
DR GO: GO:0005216; F:Ion channel activity; IEA.
DR InterPro: IPR010526; Na_trans_assoc.
DR Pfam: PF06512; Na_trans_assoc; 1.
DR Ion channel.
KW NON_TER 1
FT NON_TER 276
SQ

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SQ SEQUENCE 276 AA; 30906 MW; 5660535780CA79B3 CRC64;
Query Match 100.0%; Score 28; DB 2; Length 276;
Best Local Similarity 100.0%; Pred. No. 5e-18;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CLTVFMVWVIGNLVNLFLALLSSSF 28
Db 15 CLTVFMVWVIGNLVNLFLALLSSSF 42

RESULT 3
Q80U0U PRELIMINARY; PRT; 500 AA.
ID Q80U0U;
AC Q80U0U;
DT 01-JUN-2003 (TREMBlrel. 24, Created)
DT 01-JUN-2003 (TREMBlrel. 24, last sequence update)
DE 01-MAR-2004 (TREMBlrel. 26, last annotation update)
DE Sodium channel 5 (Fragment).
OS Ictalurus punctatus (Channel catfish).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Siluriformes;
OC Ictaluridae; Ictalurus.
OX NCBI_TaxID=7998;
RN [1]
RP SEQUENCE FROM N.A.
RA Lu Y., Lopreato G.F., Zakon H.H.;
RL Submitted (DEC-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL: AY204536; AAO60424.1; -.
DR GO: GO:0016021; C:Integral to membrane; IEA.
DR GO: GO:0005261; F:cation channel activity; IEA.
DR GO: GO:0006812; P:cation transport; IEA.
DR InterPro: IPR001682; Ca/Na_pore.
DR InterPro: IPR005821; Ion trans.
DR InterPro: IPR005820; M_channel_1lg.
DR InterPro: IPR010526; Na_trans_assoc.
DR Pfam: PF00520; Ion_trans_1.
DR Pfam: PF06512; Na_trans_assoc; 1.
DR Ion transport; Ionic channel; Transmembrane; Transport.
KW NON_TER 1
FT NON_TER 500
SQ SEQUENCE 500 AA; 56868 MW; C7DB33040416FE3F CRC64;

Query Match 100.0%; Score 28; DB 2; Length 500;
Best Local Similarity 100.0%; Pred. No. 8.1e-18;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CLTVFMVWVIGNLVNLFLALLSSSF 28
Db 6 CLTVFMVWVIGNLVNLFLALLSSSF 33

RESULT 4
Q8IU06 PRELIMINARY; PRT; 1981 AA.
ID Q8IU06;
AC Q8IU06;
DT 01-MAR-2003 (TREMBlrel. 23, Created)
DT 01-MAR-2003 (TREMBlrel. 23, last sequence update)
DE 01-MAR-2004 (TREMBlrel. 26, last annotation update)
DE Voltage-gated sodium channel alpha 1 subunit.
GN Name=SCN1A;
OS Homo sapiens (human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX TISSUE=Normal brain;
Ouchida M., Ohmori I.;
RA Submitted (DEC-2002) to the EMBL/GenBank/DBJ databases.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
CC -1- SIMILARITY: Belongs to the sodium channel family.
DR EMBL: AB098335; BAC45228.1; -.
SQ

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DR HSSP; P04775; 1BY.
DR GO; GO:0016021; C:Integral to membrane; IEA.
DR GO; GO:0001518; C:voltage-gated sodium channel complex; IEA.
DR GO; GO:0005261; F:cation channel activity; IEA.
DR GO; GO:0005248; F:voltage-gated sodium channel activity; IEA.
DR GO; GO:0006812; F:cation transport; IEA.
DR GO; GO:0006814; F:sodium ion transport; IEA.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR002111; Cat channel_TrpL.
DR InterPro; IPR005821; Ion channel_TrpL.
DR InterPro; IPR000048; Ion_trans.
DR InterPro; IPR005820; M-channel_nlg.
DR InterPro; IPR001696; Na_channel.
DR InterPro; IPR008051; Na_channel1.
DR InterPro; IPR010526; Na_trans_assoc.
DR InterPro; IPR000100; Ribonuclease_P.
DR Pfam; PF00520; Ion_trans_4.
DR Pfam; PF00612; IQ_1.
DR Pfam; PF06512; Na_trans_assoc_1.
DR PRINTS; PRO0170; NACHANNEL.
DR PRINTS; PRO1664; NACHANNEL1.
DR SMART; SMO0015; IQ_1.
DR PROSITE; PS00648; RIBONUCLEASE_P; UNKNOWN_1.
DR Ion transport; Ionic channel; Sodium channel; Transmembrane;
DR Transport; Voltage-gated channel.
DR SEQUENCE 1981 AA; 226201 MW; B1D6946D6491B7AD CRC64;

Query March 100.0%; Score 28; DB 2; Length 1981;
Best Local Similarity 100.0%; Pred. No. 2.5e-17;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Cy 1 CLTVFMVMVIGNLVNLFLALLISSF 28
Db 940 CLTVFMVMVIGNLVNLFLALLISSF 967

RESULT 5
CIN2 HUMAN STANDARD; PRT; 2005 AA.
AC Q99250; Q14472; Q9BZC9; Q9BZD0;
DT 01-JUN-1994 (Rel. 29, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 05-JUL-2004 (Rel. 44, Last annotation update)
DE Sodium channel protein type II alpha subunit (Voltage-gated sodium
DE channel alpha subunit Nav1.2) (Sodium channel protein, brain II alpha
DE subunit) (HBC II).
GN Name=SCN2A; Synonyms=SCN2A2, NAC2;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A. (ISOFORM 1).
RC TISSUE=Brain;
RA MEDLINE=92390418; PubMed=1325650;
RA Ahmed C.M., Ware D.H., Lee S.C., Patren C.D., Ferrer-Montiel A.V.,
RA Schindler A.F., McPherson J.D., Wagner-McPherson C.B., Waemuth U.J.,
RA Evans G.A., Montiel M.;
RA "Primary structure, chromosomal localization, and functional
RT expression of a voltage-gated sodium channel from human brain.";
RT Proc. Natl. Acad. Sci. U.S.A. 89:8220-8224(1992).
RN [2]
RP SEQUENCE FROM N.A. (ISOFORMS 1 AND 2).
RX MEDLINE=21142400; PubMed=11245985; DOI=10.1016/S0378-1119(00)00594-1;
RA Kasai N., Fukushima K., Ueki Y., Praad S., Nosakowski J.,
RA Sugata K.-I., Sugata A., Nishizaki K., Meyer N.C., Smith R.J.H.;
RT "Genomic structures of SCN2A and SCN3A - candidate genes for deafness
RT at the DFN16 locus.";
RT Gene 264:113-122(2001).
RN [3]
RP SEQUENCE OF 1-89 FROM N.A.
RA Lu C.-M., Eicheleberger J.S., Beckman M.L., Schade S.D., Brown G.B.;
RT "Isolation of the 5'-flanking region for human brain sodium channel

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RT subtype II alpha-Subunit (SCN2A).";
RL Submitted (Apr-1998) to the EMBL/GenBank/DBJ databases.
RN [4]
RP SEQUENCE OF 1702-2005 FROM N.A.
RC TISSUE=Brain;
RX MEDLINE=92275082; PubMed=1317301;
RA Lu C.-M., Han J., Rado T.A., Brown G.B.;
RT "Differential expression of two sodium channel subtypes in human
RT brain.";
RT FEBS Lett. 303:53-58(1992).
RN [5]
RP SEQUENCE OF 1702-1772 FROM N.A.
RX MEDLINE=91110524; PubMed=1846440;
RA Han J., Lu C.-M., Brown G.B., Rado T.A.;
RT "Direct amplification of a single dissected chromosomal segment by
RT polymerase chain reaction: a human brain sodium channel gene is on
RT chromosome 2q22-q23.";
RL Proc. Natl. Acad. Sci. U.S.A. 88:335-339(1991).
CC -1- FUNCTION: Mediates the voltage-dependent sodium ion permeability
CC of excitable membranes. Assuming opened or closed conformations in
CC response to the voltage difference across the membrane, the
CC protein forms a sodium-selective channel through which Na(+) ions
CC may pass in accordance with their electrochemical gradient.
CC -1- SUBUNIT: The sodium channel consists of a large polypeptide and 2-
CC 3 smaller ones. This sequence represents a large polypeptide.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
CC -1- ALTERNATIVE PRODUCTS:
CC Event=Alternative splicing; Named isoforms=2;
CC Name=1; Synonyms=Adult, 6A;
CC IsoId=Q99250-1; Sequence=Displayed;
CC Name=2; Synonyms=Neonatal, 6N;
CC IsoId=Q99250-2; Sequence=VSP_001032;
CC -1- DOMAIN: The sequence contains 4 internal repeats, each with 5
CC hydrophobic segments (S1,S2,S3,S5,S6) and one positively charged
CC segment (S4). Segments S4 are probably the voltage-sensors and are
CC characterized by a series of positively charged amino acids at
CC every third position.
CC -1- SIMILARITY: Belongs to the sodium channel family.
CC -1- SIMILARITY: Contains 1 IQ domain.
CC
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CC
CC EMBL; M94055; AAA18895.1; -
CC EMBL; AF059683; AAC14574.1; -
CC EMBL; AF327246; AAG53413.1; -
CC EMBL; AF327226; AAG53413.1; JOINED.
CC EMBL; AF327227; AAG53413.1; JOINED.
CC EMBL; AF327228; AAG53413.1; JOINED.
CC EMBL; AF327229; AAG53413.1; JOINED.
CC EMBL; AF327230; AAG53413.1; JOINED.
CC EMBL; AF327231; AAG53413.1; JOINED.
CC EMBL; AF327232; AAG53413.1; JOINED.
CC EMBL; AF327233; AAG53413.1; JOINED.
CC EMBL; AF327234; AAG53413.1; JOINED.
CC EMBL; AF327235; AAG53413.1; JOINED.
CC EMBL; AF327236; AAG53413.1; JOINED.
CC EMBL; AF327237; AAG53413.1; JOINED.
CC EMBL; AF327238; AAG53413.1; JOINED.
CC EMBL; AF327239; AAG53413.1; JOINED.
CC EMBL; AF327240; AAG53413.1; JOINED.
CC EMBL; AF327241; AAG53413.1; JOINED.
CC EMBL; AF327242; AAG53413.1; JOINED.
CC EMBL; AF327243; AAG53413.1; JOINED.
CC EMBL; AF327244; AAG53413.1; JOINED.
CC EMBL; AF327245; AAG53413.1; JOINED.
CC EMBL; AF327246; AAG53412.1; -
CC EMBL; AF327226; AAG53412.1; JOINED.

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Query Match	Best Local Similarity	100.0%;	Score 28;	DB 1;	Length 2005;
Matches 28;	Conservative	0;	Mismatches 0;	Indels 0;	Gaps 0
FT CARBOHYD	285	285	N-linked (GlcNAc. . .)	(Potential)	.
FT CARBOHYD	291	291	N-linked (GlcNAc. . .)	(Potential)	.
FT CARBOHYD	297	297	N-linked (GlcNAc. . .)	(Potential)	.
FT CARBOHYD	303	303	N-linked (GlcNAc. . .)	(Potential)	.
FT CARBOHYD	308	308	N-linked (GlcNAc. . .)	(Potential)	.
FT CARBOHYD	340	340	N-linked (GlcNAc. . .)	(Potential)	.
FT CARBOHYD	604	604	N-linked (GlcNAc. . .)	(Potential)	.
FT CARBOHYD	824	624	N-linked (GlcNAc. . .)	(Potential)	.
FT CARBOHYD	883	883	N-linked (GlcNAc. . .)	(Potential)	.
FT CARBOHYD	1055	1055	N-linked (GlcNAc. . .)	(Potential)	.
FT CARBOHYD	1072	1072	N-linked (GlcNAc. . .)	(Potential)	.
FT CARBOHYD	1136	1136	N-linked (GlcNAc. . .)	(Potential)	.
FT CARBOHYD	1368	1368	N-linked (GlcNAc. . .)	(Potential)	.
FT CARBOHYD	1382	1382	N-linked (GlcNAc. . .)	(Potential)	.
FT CARBOHYD	1393	1393	N-linked (GlcNAc. . .)	(Potential)	.
FT CARBOHYD	1778	1778	N-linked (GlcNAc. . .)	(Potential)	.
FT VARSPLIC	209	209	D -> N (in isoform 2).	(Potential)	.
FT CONFLICT	524	524	/FTId=VSP_001032.		
			R -> L (in Ref. 1).		
Query Match	Best Local Similarity	100.0%;	Score 28;	DB 1;	Length 2005;
Matches 28;	Conservative	0;	Mismatches 0;	Indels 0;	Gaps 0
Qy	1	CLTFPMWVWVGNLVNLFALTLSSSF 28			
Db	955	CLTFPMWVWVGNLVNLFALTLSSSF 966			
RESULT 6					
CIN2_RAT					
ID	CIN2_RAT	STANDARD;	PRT;	2005 AA.	
AC	P04775;				
DT	13-AUG-1987 (Rel. 05. Created)				
DT	13-AUG-1987 (Rel. 05. Last sequence update)				
DT	01-OCT-2004 (Rel. 45. Last annotation update)				
DE	Sodium channel protein type II alpha subunit (Voltage-gated sodium channel alpha subunit Nav1.2) (Sodium channel protein, brain II alpha subunit).				
DE	Name=Scn2a;				
GN	Rattus norvegicus (Rat).				
OS	Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;				
OC	Mammalia; Euteleostomi; Rodentia; Sciurognathi; Muridae; Rattus.				
OX	NCBI_TaxID=10116;				
RA	NCBI_TaxID=10116;				
RP	SEQUENCE FROM N.A.				
RX	MEDLINE=86146901; Pubmed=3754035;				
RA	Noda M., Ikeda T., Kayano T., Suzuki H., Takeshima H., Kuraaki M.,				
RA	Takahashi H., Numa S.;				
RT	"Existence of distinct sodium channel messenger RNAs in rat brain.";				
RL	Nature 320:188-192(1986).				
CC	-I- FUNCTION: Mediates the voltage-dependent sodium ion permeability of excitable membranes. Assuming opened or closed conformations in response to the voltage difference across the membrane, the protein forms a sodium-selective channel through which Na(+) ions may pass in accordance with their electrochemical gradient.				
CC	-I- SUBUNIT: The sodium channel consists of a large polypeptide and 2-3 smaller ones. This sequence represents a large polypeptide.				
CC	-I- SUBCELLULAR LOCATION: Integral membrane protein.				
CC	-I- DOMAIN: The sequence contains 4 internal repeats, each with 5 hydrophobic segments (S1,S2,S3,S5,S6) and one positively charged segment (S4). Segments S4 are probably the voltage-sensors and are characterized by a series of positively charged amino acids at every third position.				
CC	-I- SIMILARITY: Belongs to the sodium channel family.				
CC	-I- SIMILARITY: Contains 1 IQ domain.				
CC	-----				
CC	This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities remains prohibited.				

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CC EMBL; X03639; CAA27287.1; -.
DR PDB; 1BY; NMR; A=1474-1526.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR002111; Cat_channel_TrypL.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IQ_region.
DR InterPro; IPR005820; M-channel_nlg.
DR InterPro; IPR001696; Na_channel.
DR InterPro; IPR010526; Na_trans_assoc.
DR Pfam; PF00520; Ion_trans_4.
DR Pfam; PF06512; IQ_1.
DR Pfam; PF06512; Na_trans_assoc_1.
DR PRINTS; PR00170; NACHANNEL.
DR PROSITE; PS0096; IQ_1.
KW 3D-structure; Glycoprotein; Ion transport; Ionic channel;
KW Multigene family; Repeat; Sodium channel; Transmembrane;
KW Voltage-gated channel.
FT REPEAT 111 456 I.
FT REPEAT 741 1013 II.
FT REPEAT 1190 1504 III.
FT REPEAT 1513 1811 IV.
FT TRANSMEM 125 148 S1 of repeat I.
FT TRANSMEM 157 176 S2 of repeat I.
FT TRANSMEM 190 208 S3 of repeat I.
FT TRANSMEM 215 234 S4 of repeat I.
FT TRANSMEM 251 274 S5 of repeat I.
FT TRANSMEM 402 427 S6 of repeat I.
FT TRANSMEM 754 778 S1 of repeat II.
FT TRANSMEM 822 841 S2 of repeat II.
FT TRANSMEM 884 904 S3 of repeat II.
FT TRANSMEM 958 983 S4 of repeat II.
FT TRANSMEM 1204 1227 S5 of repeat II.
FT TRANSMEM 1241 1266 S6 of repeat II.
FT TRANSMEM 1273 1294 S1 of repeat III.
FT TRANSMEM 1299 1320 S2 of repeat III.
FT TRANSMEM 1340 1367 S3 of repeat III.
FT TRANSMEM 1447 1473 S4 of repeat III.
FT TRANSMEM 1527 1550 S5 of repeat III.
FT TRANSMEM 1562 1585 S6 of repeat III.
FT TRANSMEM 1592 1615 S1 of repeat IV.
FT TRANSMEM 1626 1647 S2 of repeat IV.
FT TRANSMEM 1663 1685 S3 of repeat IV.
FT TRANSMEM 1752 1776 S4 of repeat IV.
FT DOMAIN 1905 1934 S5 of repeat IV.
FT CARBOHYD 212 212 IQ.
FT CARBOHYD 285 285 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 291 291 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 297 297 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 303 303 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 308 308 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 340 340 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 604 604 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 624 624 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 883 883 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 1055 1055 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 1072 1072 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 1136 1136 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 1368 1368 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 1382 1382 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 1393 1393 N-linked (GlcNAc...) (Potential).
SQ SEQUENCE 2005 AA; 227872 MW; 861BE583D79F8324 CRC64;

Query Match 100.0%; Score 28; DB 1; Length 2005;

Best Local Similarity 100.0%; Pred. No. 2.Se-17; Indels 0; Gaps 0;

Db 1 CLTVFMVMVIGNLVNLFLALLLSF 28
959 CLTVFMVMVIGNLVNLFLALLLSF 986

RESULT 7

O9YGN7

ID O9YGN7

AC O9YGN7

DT 01-MAY-1999 (TRENBLrel. 10, Created)

DT 01-MAY-1999 (TRENBLrel. 10, Last sequence update)

DT 01-MAR-2004 (TRENBLrel. 26, Last annotation update)

DE Voltage-dependent sodium channel.

OS Cynops pyrrhogaster (Japanese common newt).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Amphibia; Batrachia; Caudata; Salamandridae; Cynops.

OX NCBI_Taxid=8330;

RN [1]

RP SEQUENCE FROM N.A.

RC TISSUE=Retina;

RA Hirota K., Kaneko Y., Matsumoto G., Hanyu Y.;

RL Submitted (JAN-1999) to the EMBL/GenBank/DBJ databases.

CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).

CC -1- SIMILARITY: Belongs to the sodium channel family.

DR EMBL; AF123593; AAD17315.1; -.

DR HSSP; P04775; 1BY.

DR GO; GO:0016021; C:Integral to membrane; IEA.

DR GO; GO:0001518; C:Voltage-gated sodium channel complex; IEA.

DR GO; GO:0005261; F:cation channel activity; IEA.

DR GO; GO:0005248; F:voltage-gated sodium channel activity; IEA.

DR GO; GO:0006812; P:cation transport; IEA.

DR GO; GO:0006814; P:sodium ion transport; IEA.

DR InterPro; IPR001682; Ca/Na_pore.

DR InterPro; IPR002111; Cat_channel_TrypL.

DR InterPro; IPR005821; Ion_trans.

DR InterPro; IPR000048; IQ_region.

DR InterPro; IPR005820; M-channel_nlg.

DR InterPro; IPR01696; Na_channel.

DR Pfam; PF00520; Ion_trans_4.

DR Pfam; PF06512; IQ_1.

DR Pfam; PF06512; Na_trans_assoc_1.

DR PRINTS; PR00170; NACHANNEL.

DR SMART; SM00015; IQ_1.

KW Ion transport; Ionic channel; Sodium channel; Transmembrane;

KW Transport; Voltage-gated channel.

SQ SEQUENCE 2007 AA; 228398 MW; 013B9B9C9C294C9 CRC64;

Query Match 100.0%; Score 28; DB 2; Length 2007;
Best Local Similarity 100.0%; Pred. No. 2.Se-17;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CLTVFMVMVIGNLVNLFLALLLSF 28
Db 957 CLTVFMVMVIGNLVNLFLALLLSF 984

RESULT 8

CINI_HUMAN

ID CINI_HUMAN

AC P35498; Q16172; Q961A3; Q9C008;

DT 01-JUN-1994 (Rel. 29, Created)

DT 16-OCT-2001 (Rel. 40, Last sequence update)

DT 05-JUL-2004 (Rel. 44, Last annotation update)

DE Sodium channel protein type I alpha subunit (Voltage-gated sodium

DE channel alpha subunit Nav1.1) (Sodium channel protein, Brain I alpha

DE subunit).

DE Name=SCN1A; Synonym=SCN1, NAC1;

OS Homo sapiens (Human).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

OX NCBI_Taxid=9606;

RN [1]

RP SEQUENCE FROM N.A. (ISOFORM 1), AND VARIANTS GEF5+2 MET-875 AND

RP RP HIS-1648.

RX MEDLINE=20206553; PubMed=10742094;

RA Escayg A., MacDonald B.T., Weiler M.H., Baulac S., Huberfeld G.,
 RA An-Gourfinkel I., Brice A., Ieuearn E., Moulard B., Chaigne D.,
 RA Buresi C., Malafosse A.;
 RT "Mutations of SCN1A, encoding a neuronal sodium channel, in two
 RT families with GERS+2.";
 RL Nat. Genet. 24:343-345(2000).
 RN (12)
 RP SEQUENCE FROM N.A. (ISOFORM 2).
 RA Jeong S.-Y., Goto J., Kanazawa I.;
 RT "Cloning of cDNA for human voltage-gated sodium channel alpha subunit,
 RT SCN1A.";
 RL Submitted (JAN-2000) to the EMBL/GenBank/DBJ databases.
 RN (13)
 RP SEQUENCE FROM N.A. (ISOFORM 2).
 RC TISSUE=Brain;
 RA Sugawara T., Mazaki E.M., Yamakawa K.;
 RT "Homo sapiens neuronal voltage-gated sodium channel type I (Nav1.1)
 RT mRNA.";
 RL Submitted (JUL-2001) to the EMBL/GenBank/DBJ databases.
 RN (14)
 RP SEQUENCE FROM N.A. (ISOFORMS 1 AND 2), AND VARIANT THR-1067.
 RA Ouchida M., Ohmori I.;
 RT "isoforms of human sodium channel SCN1A gene.";
 RL Submitted (OCT-2002) to the EMBL/GenBank/DBJ databases.
 RN (15)
 RP SEQUENCE OF 1335-1428 FROM N.A.
 RX MEDLINE=94340991; PubMed=8062593;
 RA Malo M.S., Blanchard B.J., Andresen J.M., Srivastava K., Chen X.N.,
 RA Li X., Jabs E.W., Korenberg J.R., Ingiram V.M.;
 RT "Localization of a putative human brain sodium channel gene (SCN1A) to
 RT chromosome band 2q24.";
 RL Cytogenet. Cell Genet. 67:178-186(1994).
 RN (16)
 RP SEQUENCE OF 1518-1940 FROM N.A.
 RC TISSUE=Brain;
 RX MEDLINE=92275082; PubMed=1117301;
 RA Lu C.-W., Han J., Rado T.A., Brown G.B.;
 RT "Differential expression of two sodium channel subtypes in human
 RT brain.";
 RL FEBS Lett. 303:53-58(1992).
 RN (17)
 RP VARIANTS GERS+2 VAL-188, LEU-1353 AND MET-1656.
 RX MEDLINE=21152274; PubMed=11254444;
 RA Wallace R.H., Scheffer I.E., Barnett S., Richards M., Dibbens L.,
 RA Desai R.R., Iernan-Sagle T., Lev D., Mazarib A., Brand N.,
 RA Ben-Zeev B., Golthman I., Singh R., Kremmidiotis G., Gardner A.,
 RA Suberland G.R., George A.L. Jr., Mulley J.C., Berkovic S.F.;
 RT "Neuronal sodium-channel alpha-subunit mutations in generalized
 RT epilepsy with febrile seizures plus.";
 RL Am. J. Hum. Genet. 68:859-865(2001).
 RN (18)
 RP VARIANTS GERS+2 ARG-1204.
 RX MEDLINE=21152275; PubMed=11254445;
 RA Escayg A., Heils A., MacDonald B.T., Haug K., Sander T., Meisler M.H.;
 RT "A novel SCN1A mutation associated with generalized epilepsy with
 RT febrile seizures plus -- and prevalence of variants in patients with
 RT epilepsy.";
 RL Am. J. Hum. Genet. 68:866-873(2001).
 RN (19)
 RP VARIANT SWEI PHE-986.
 RX MEDLINE=21257503; PubMed=11359211;
 RA Cies L., Del-Favero J., Culemans B., Lagae L., Van Broeckhoven C.,
 RA De Jonghe P.;
 RT "De novo mutations in the sodium-channel gene SCN1A cause severe
 RT myoclonic epilepsy of infancy.";
 RL Am. J. Hum. Genet. 68:1327-1332(2001).
 RN (10)
 RP VARIANT GERS+2 THR-1270.
 RX MEDLINE=21630138; PubMed=11756608;
 RA Abou-Khalil B., Ge O., Desai R., Ryther R., Bazzyk A., Bailey R.,
 RA Haines J.L., Sutcliffe J.S., George A.L. Jr.;
 RT "Partial and generalized epilepsy with febrile seizures plus and a
 RT novel SCN1A mutation.";

RL Neurology 57:2265-2272(2001).
 CC -1- FUNCTION: Mediates the voltage-dependent sodium ion permeability
 CC of excitable membranes. Assuming opened or closed conformations in
 CC response to the voltage difference across the membrane, the
 CC protein forms a sodium-selective channel through which Na(+) ions
 CC may pass in accordance with their electrochemical gradient.
 CC -1- SUBUNIT: The sodium channel consists of a large polypeptide and 2-
 CC 3 smaller ones. This sequence represents a large polypeptide.
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
 CC -1- ALTERNATIVE PRODUCTS:
 CC Event=Alternative splicing; Named isoforms=2;
 CC Name=1;
 CC IsoId=P35498-1; Sequence=Displayed;
 CC Name=2;
 CC IsoId=P35498-2; Sequence=VSP_001011;
 CC Note=No experimental confirmation available;
 CC -1- DOMAIN: The sequence contains 4 internal repeats, each with 5
 CC hydrophobic segments (S1,S2,S3,S5,S6) and one positively charged
 CC segment (S4). Segments S4 are probably the voltage-sensors and are
 CC characterized by a series of positively charged amino acids at
 CC every third position.
 CC -1- DISEASE: Defects in SCN1A are the cause of generalized epilepsy
 CC with febrile seizures plus type 2 (GERS+2) [MIM:604233]. This
 CC autosomal dominant disorder is characterized by febrile seizures
 CC in children and afebrile seizures in adults. Penetrance is
 CC incomplete and a large intrafamilial variability of the phenotype
 CC is observed.
 CC -1- DISEASE: Defects in SCN1A are a cause of severe myoclonic epilepsy
 CC in infancy (SWEI) [MIM:607208], a severe form of generalized
 CC epilepsy with febrile seizures. SWEI is a rare disorder
 CC characterized by normal development before onset, seizures
 CC beginning in the first year of life in the form of generalized or
 CC unilateral febrile clonic seizures, secondary appearance of
 CC myoclonic seizures, and occasionally partial seizures. It is
 CC associated with ataxia, slowed psychomotor development, and mental
 CC decline.
 CC -1- SIMILARITY: Belongs to the sodium channel family.
 CC -1- SIMILARITY: Contains 1 IQ domain.
 CC -----
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 CC -----
 CC EMBL; AF225985; AAK00217.1; -;
 CC EMBL; AY043484; AAK95360.1; -;
 CC EMBL; AB093548; BAC21101.1; -;
 CC EMBL; AB093549; BAC21102.1; -;
 CC EMBL; S71446; AAB31605.1; -;
 CC EMBL; X65362; CAA46439.1; -;
 CC EMBL; M91803; -; NOT_ANNOTATED_CDS.
 CC PIR; I52964; I52964.
 CC PIR; S29184; S29184.
 CC HSSP; P04775; 1BY7.
 CC Genew; HGNC:10585; SCN1A.
 CC MIM; 182389; -;
 CC MIM; 604233; -;
 CC MIM; 607208; -;
 CC GO; GO:0016021; C:integral to membrane; NAS.
 CC GO; GO:0005248; F:voltage-gated sodium channel activity; NAS.
 CC GO; GO:0006814; P:sodium ion transport; NAS.
 CC InterPro; IPR001682; Ca/Na_pore.
 CC InterPro; IPR002111; Cat_channel_TrypL.
 CC InterPro; IPR005821; Ion_trans.
 CC InterPro; IPR000048; IQ_region.
 CC InterPro; IPR005820; M+channel_nlg.
 CC InterPro; IPR001696; Na_channel.
 CC InterPro; IPR008051; Na_channel1.
 CC InterPro; IPR010526; Na_trans_assoc.
 CC Pfam; Pf00520; Ion_trans; 4.

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DR Pfam; PF00612; IQ; 1.
DR Pfam; PF06512; Na_trans_assoc; 1.
DR PRINTS; PRO0170; NACHANNEL.
DR PRINTS; PRO1664; NACHANNEL.
DR PROSITE; PS0096; IQ; FALSE_NEG.
KW Alternative splicing; Disease mutation; Epilepsy; Glycoprotein;
KW Ion transport; Ionic channel; Multigene family; Polymorphism; Repeat;
KW Sodium channel; Transmembrane; Voltage-gated channel.
FT REPEAT 110 454
FT REPEAT 750 1022
FT REPEAT 1200 1514
FT REPEAT 1523 1821
FT TRANSMEM 124 147
FT TRANSMEM 156 175
FT TRANSMEM 189 207
FT TRANSMEM 214 233
FT TRANSMEM 250 273
FT TRANSMEM 400 425
FT TRANSMEM 763 787
FT TRANSMEM 799 822
FT TRANSMEM 831 850
FT TRANSMEM 857 876
FT TRANSMEM 893 913
FT TRANSMEM 967 992
FT TRANSMEM 1214 1237
FT TRANSMEM 1251 1276
FT TRANSMEM 1283 1304
FT TRANSMEM 1309 1330
FT TRANSMEM 1350 1377
FT TRANSMEM 1457 1483
FT TRANSMEM 1537 1560
FT TRANSMEM 1572 1595
FT TRANSMEM 1602 1625
FT TRANSMEM 1636 1657
FT TRANSMEM 1673 1695
FT TRANSMEM 1762 1786
FT CARBOHYD 211 211

Query Match 100.0%; Score 28; DB 1; Length 2009;
Best Local Similarity 100.0%; Pred. No. 2.5e-17;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CLTVFMVWVIGNLVNLFLLALLSSF 28
Db 968 CLTVFMVWVIGNLVNLFLLALLSSF 995

RESULT 9
Q800U4 ID Q800U4 PRELIMINARY; PRT; 472 AA.
AC Q800U4.
DT 01-JUN-2003 (TREMBLrel. 24, Created)
DT 01-JUN-2003 (TREMBLrel. 24, Last sequence update)
DT 01-MAR-2004 (TREMBLrel. 26, Last annotation update)
DE Sodium channel 1 (Fragment).
OS Ictalurus punctatus (Channel catfish).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Siluriformes;
OC Ictaluridae; Ictalurus.
OX NCBI_TaxID=7998;
RN [1]
RP SEQUENCE FROM N.A.
RA Lu Y., Lopreato G.F., Zakon H.H.;
RL Submitted (DEC-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; AY204532; AA060420.1; -.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0005261; F:cation channel activity; IEA.
DR GO; GO:0006812; P:cation transport; IEA.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR005820; M-channel_nlg.
DR InterPro; IPR010526; Na_trans_assoc.
DR Pfam; PF00520; Ion_trans; 1.

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DR Pfam; PF06512; Na_trans_assoc; 1.
KW Ion transport; Ionic channel; Transmembrane; Transport.
FT NON_TER 1
FT NON_TER 472
FT NON_TER 472
SQ SEQUENCE 472 AA; 53917 MW; 3DD68F4EAB484FAB CRC64;

Query Match 89.3%; Score 25; DB 2; Length 472;
Best Local Similarity 100.0%; Pred. No. 4.8e-15;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 4 VPMWVWVIGNLVNLFLLALLSSF 28
Db 9 VPMWVWVIGNLVNLFLLALLSSF 33

RESULT 10
Q902Z6 ID Q902Z6 PRELIMINARY; PRT; 743 AA.
AC Q902Z6.
DT 01-DEC-2001 (TREMBLrel. 19, Created)
DT 01-DEC-2001 (TREMBLrel. 19, Last sequence update)
DT 01-MAR-2004 (TREMBLrel. 26, Last annotation update)
DE Sodium channel 5 (Fragment).
OS Sternopygus macrurus.
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Gymnotiformes;
OC Sternopygidae; Sternopygus.
OX NCBI_TaxID=77841;
RN [1]
RP SEQUENCE FROM N.A.
RA MEDLINE=21310016; PubMed=11416226;
RA Lopreato G.F., Lu Y., Southwell A., Atkinson N.S., Hillis D.M.,
RA Wilcox T.P., Zakon H.H.;
RT "Evolution and divergence of sodium channel genes in vertebrates."
RL Proc. Natl. Acad. Sci. U.S.A. 98:7588-7592(2001).
DR EMBL; AF378143; AAK55441.1; -.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0005261; F:cation channel activity; IEA.
DR GO; GO:0006812; P:cation transport; IEA.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR002111; Cat_channel_TrpL.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR005820; M-channel_nlg.
DR InterPro; IPR010526; Na_trans_assoc.
DR Pfam; PF00520; Ion_trans; 2.
DR Pfam; PF06512; Na_trans_assoc; 1.
KW Ion transport; Ionic channel; Transmembrane; Transport.
FT NON_TER 1
FT NON_TER 743
FT NON_TER 743
SQ SEQUENCE 743 AA; 84781 MW; F2429665544CABOC CRC64;

Query Match 89.3%; Score 25; DB 2; Length 743;
Best Local Similarity 100.0%; Pred. No. 7e-15;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 4 VPMWVWVIGNLVNLFLLALLSSF 28
Db 182 VPMWVWVIGNLVNLFLLALLSSF 206

RESULT 11
Q902Z8 ID Q902Z8 PRELIMINARY; PRT; 751 AA.
AC Q902Z8.
DT 01-DEC-2001 (TREMBLrel. 19, Created)
DT 01-DEC-2001 (TREMBLrel. 19, Last sequence update)
DT 01-MAR-2004 (TREMBLrel. 26, Last annotation update)
DE Sodium channel 3 (Fragment).
OS Sternopygus macrurus.
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Gymnotiformes;
OC Sternopygidae; Sternopygus.
OX NCBI_TaxID=77841;

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RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=21310016; PubMed=11416226;
RA Lopezato G.F., Lu Y., Southwell A., Atkinson N.S., Hillis D.M.,
RT "Evolution and divergence of sodium channel genes in vertebrates.";
RL Proc. Natl. Acad. Sci. U.S.A. 98:7588-7592(2001).
DR EMBL; AF378141; AKS5439.1; -.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0005261; P:cation channel activity; IEA.
DR GO; GO:0006812; P:cation transport; IEA.
DR InterPro; IPR001682; Ca/Na pore.
DR InterPro; IPR002111; Cat_channel_TrpL.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR005820; M+channel_nlg.
DR Pfam; PF00520; Ion_trans_2.
DR Pfam; PF06512; Na_trans_assoc; 1.
KM Ion transport; Ionic channel; Transmembrane; Transport.
FT NON_TER
FT NON_TER
SQ SEQUENCE 751 AA; 84598 MW; CBFE6162E30A76FC CRC64;

Query Match
Best Local Similarity 100.0%; Pred. No. 7e-15; Length 751;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 4 VFMWVVGIVGNLVVNLFLALLISSF 28
Db 182 VFMWVVGIVGNLVVNLFLALLISSF 206

RESULT 12
ID Q90519 PRELIMINARY; PRT; 1717 AA.
AC Q90519;
DT 01-NOV-1996 (TREMBLrel. 01, Created)
DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
DT 01-MAR-2004 (TREMBLrel. 26, Last annotation update)
DS Sodium channel alpha subunit (Fragment).
OS Fugu rubripes (Japanese pufferfish) (Takifugu rubripes).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
OC Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes;
OC Tetraodontidae; Tetraodontidae; Takifugu.
OX NCBI_TaxID=31033;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Brain;
RA Nakazawa A.;
RL Submitted (AUG-1994) to the EMBL/GenBank/DBJ databases.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
CC -1- SIMILARITY: Belongs to the sodium channel family.
DR EMBL; D37977; BAA07195.1; -.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0001518; C:voltage-gated sodium channel complex; IEA.
DR GO; GO:0005261; F:cation channel activity; IEA.
DR GO; GO:0005248; F:voltage-gated sodium channel activity; IEA.
DR GO; GO:0006812; P:cation transport; IEA.
DR GO; GO:0006814; P:sodium ion transport; IEA.
DR InterPro; IPR001682; Ca/Na pore.
DR InterPro; IPR002111; Cat_channel_TrpL.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IQ_region.
DR InterPro; IPR005820; M+channel_nlg.
DR InterPro; IPR001696; Na_channel.
DR InterPro; IPR008054; Na_channel8.
DR InterPro; IPR010526; Na_trans_assoc.
DR Pfam; PF00520; Ion_trans_4.
DR Pfam; PF06512; Na_trans_assoc; 1.
DR PRINTS; PR00170; NACHANNEL.
DR PRINTS; PR01667; NACHANNEL8.

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DR SMART; SM00015; IQ; 1.
DR PROSITE; PS50096; IQ; 1.
KM Ion transport; Ionic channel; Sodium channel; Transmembrane;
KW Transport; Voltage-gated channel.
FT NON_TER
FT NON_TER
SQ SEQUENCE 1717 AA; 195413 MW; 067D566B3FA41C8D CRC64;

Query Match
Best Local Similarity 100.0%; Pred. No. 1.4e-14; Length 1717;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 4 VFMWVVGIVGNLVVNLFLALLISSF 28
Db 721 VFMWVVGIVGNLVVNLFLALLISSF 745

RESULT 13
ID Q9IBF1 PRELIMINARY; PRT; 1880 AA.
AC Q9IBF1;
DT 01-OCT-2000 (TREMBLrel. 15, Created)
DT 01-OCT-2000 (TREMBLrel. 15, Last sequence update)
DT 01-MAR-2004 (TREMBLrel. 26, Last annotation update)
DS Voltage-gated sodium channel.
OS Takifugu pardalis (puffer).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
OC Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes;
OC Tetraodontidae; Tetraodontidae; Takifugu.
OX NCBI_TaxID=98921;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Skeletal muscle;
RX MEDLINE=20090650; PubMed=10623632;
RA Yonezu-Yamashita M., Nishimori K., Nitanai Y., Isejima M., Sugimoto A.,
RA Yasumoto T.;
RL "Binding properties of 3H-PbTx-3 and 3H-saxitoxin to brain membranes
and to skeletal muscle membranes of puffer fish Fugu pardalis, and the
primary structure of a voltage-gated Na+ channel alpha-subunit (FNNa1)
from skeletal muscle of F. pardalis.";
RL Biochem. Biophys. Res. Commun. 267:403-412(2000).
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
CC -1- SIMILARITY: Belongs to the sodium channel family.
DR EMBL; AB030482; BAA90398.1; -.
DR HSSP; P04775; IBY.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0001518; C:voltage-gated sodium channel complex; IEA.
DR GO; GO:0005261; F:cation channel activity; IEA.
DR GO; GO:0005248; F:voltage-gated sodium channel activity; IEA.
DR GO; GO:0006812; P:cation transport; IEA.
DR GO; GO:0006814; P:sodium ion transport; IEA.
DR InterPro; IPR001682; Ca/Na pore.
DR InterPro; IPR002111; Cat_channel_TrpL.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IQ_region.
DR InterPro; IPR005820; M+channel_nlg.
DR InterPro; IPR001696; Na_channel.
DR InterPro; IPR010526; Na_trans_assoc.
DR Pfam; PF00520; Ion_trans_4.
DR Pfam; PF06512; Na_trans_assoc; 1.
DR PRINTS; PR00170; NACHANNEL.
DR PRINTS; PS50096; IQ; 1.
KM Ion transport; Ionic channel; Sodium channel; Transmembrane;
KW Transport; Voltage-gated channel.
SQ SEQUENCE 1880 AA; 212084 MW; 406483C63D3E02 CRC64;

Query Match
Best Local Similarity 100.0%; Pred. No. 1.5e-14; Length 1880;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 4 VFMWVVGIVGNLVVNLFLALLISSF 28

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DB 803 VFMWVIGNLVNLFLALLSSP 827

RESULT 14

ID Q9DF53 PRELIMINARY; PRT; 1949 AA.

AC Q9DF53; 01-MAR-2001 (TREMBlrel. 16, Created)

DT 01-MAR-2001 (TREMBlrel. 16, Last sequence update)

DT 01-MAR-2004 (TREMBlrel. 26, Last annotation update)

DE Sodium channel protein Scn8a.

GN Name=Scn8a;

OS Brachydanio rerio (Zebrafish) (Danio rerio).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;

OC Cyprinidae; Danio.

OC NCBI_TaxId=7955;

RN (1)

RP SEQUENCE FROM N.A.

RC STRAIN=oregon;

RA Tsai C.-W., Tseng J.-J., Horng J.-F., Mu J.-L., Tsay H.-J.;

RL Submitted (AUG-2000) to the EMBL/GenBank/DBJ databases.

CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).

CC -1- SIMILARITY: Belongs to the sodium channel family.

DR EMBL; AF297658; AAG18440.1; -.

DR HSSP; P04775; 1BYX.

DR ZFIN; ZDB-GENE-000828-1; scn8a.

DR GO; GO:0016021; C:integral to membrane; IEA.

DR GO; GO:0001518; C:voltage-gated sodium channel complex; IEA.

DR GO; GO:0005261; F:cation channel activity; IEA.

DR GO; GO:0005248; F:voltage-gated sodium channel activity; IEA.

DR GO; GO:0006812; P:cation transport; IEA.

DR GO; GO:0006814; P:sodium ion transport; IEA.

DR InterPro; IPR001682; Ca/Na_pore.

DR InterPro; IPR002111; Cat_channel_TrypL.

DR InterPro; IPR005821; Ion_trans.

DR InterPro; IPR000048; IQ_region.

DR InterPro; IPR005820; M+channel_nlg.

DR InterPro; IPR001696; Na_channel.

DR InterPro; IPR008054; Na_channel18.

DR InterPro; IPR010526; Na_trans_assoc.

DR Pfam; PF00520; Ion_trans_4.

DR Pfam; PF06512; IQ_1.

DR Pfam; PF06512; Na_trans_assoc; 1.

DR PRINTS; PR00170; NACHANNEL.

DR PRINTS; PR01667; NACHANNEL8.

DR SMART; SM00015; IQ; 1.

DR Ion transport; Ionic channel; Sodium channel; Transmembrane;

KW Transport; Voltage-gated channel.

SO SEQUENCE 1949 AA; 221760 MW; 6BCA69664BDC7BC3 CRC64;

Query Match 89.3%; Score 25; DB 2; Length 1949;

Best Local Similarity 100.0%; Pred. No. 1.5e-14;

Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 VFMWVIGNLVNLFLALLSSP 28

DB 925 VFMWVIGNLVNLFLALLSSP 949

RESULT 15

ID Q63541 PRELIMINARY; PRT; 1976 AA.

AC Q63541; 01-NOV-1996 (TREMBlrel. 01, Created)

DT 01-NOV-1996 (TREMBlrel. 01, Last sequence update)

DT 01-MAR-2004 (TREMBlrel. 26, Last annotation update)

DE Sodium channel protein 6.

GN Name=SCN6;

OS Rattus norvegicus (Rat).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.

OC NCBI_TaxId=10116;

RN (1)

RP SEQUENCE FROM N.A.

RC STRAIN=Sprague-Dawley; TISSUE=Brain;

RX MEDLINE=95271284; PubMed=7751906;

RA Schaller K.L., Kizemien D.M., Yarowsky P.J., Krueger B.K.,

RA Caldwell J.H.;

RT "A novel, abundant sodium channel expressed in neurons and glia.";

RL J. Neurosci. 15:3231-3242 (1995).

CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).

CC -1- SIMILARITY: Belongs to the sodium channel family.

DR EMBL; U39018; AAC42059.1; -.

DR PIR; I56555; I56555.

DR HSSP; P04775; 1BYX.

DR GO; GO:0016021; C:integral to membrane; IEA.

DR GO; GO:0001518; C:voltage-gated sodium channel complex; IEA.

DR GO; GO:0005261; F:cation channel activity; IEA.

DR GO; GO:0005248; F:voltage-gated sodium channel activity; IEA.

DR GO; GO:0006812; P:cation transport; IEA.

DR GO; GO:0006814; P:sodium ion transport; IEA.

DR InterPro; IPR001682; Ca/Na_pore.

DR InterPro; IPR002111; Cat_channel_TrypL.

DR InterPro; IPR000183; Decarboxylase2.

DR InterPro; IPR005821; Ion_trans.

DR InterPro; IPR000048; IQ_region.

DR InterPro; IPR005820; M+channel_nlg.

DR InterPro; IPR001696; Na_channel18.

DR InterPro; IPR008054; Na_trans_assoc.

DR Pfam; PF00520; Ion_trans_4.

DR Pfam; PF00612; IQ_1.

DR Pfam; PF06512; Na_trans_assoc; 1.

DR PRINTS; PR00170; NACHANNEL.

DR PRINTS; PR01667; NACHANNEL8.

DR SMART; SM00015; IQ; 1.

DR PROSITE; PS50096; IQ; 1.

DR PROSITE; PS00878; ODR DC 2.1; UNKNOWN 1.

KW Ion transport; Ionic channel; Sodium channel; Transmembrane;

KW Transport; Voltage-gated channel.

SO SEQUENCE 1976 AA; 225227 MW; B6949327A47FA88A CRC64;

Query Match 89.3%; Score 25; DB 2; Length 1976;

Best Local Similarity 100.0%; Pred. No. 1.5e-14;

Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 VFMWVIGNLVNLFLALLSSP 28

DB 952 VFMWVIGNLVNLFLALLSSP 976

RESULT 16

ID C1N8_MOUSE STANDARD; PRT; 1978 AA.

AC Q9WTU3; Q60828; Q60858; Q62449;

DT 10-OCT-2003 (Rel. 42, Created)

DT 10-OCT-2003 (Rel. 42, Last sequence update)

DT 05-JUL-2004 (Rel. 44, Last annotation update)

DE Sodium channel protein type VIII alpha subunit (Voltage-gated sodium channel alpha subunit Nav1.6).

GN Name=Scn8a; Synonyms=Nbnal1;

OS Mus musculus (Mouse).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

OC NCBI_TaxId=10090;

RN (1)

RP SEQUENCE FROM N.A. (ISOFORMS 1 AND 3).

RC STRAIN=C57BL/6J;

RX MEDLINE=99047535; PubMed=9828131;

RA Plummer N.W., Galt J., Jones J.M., Burgess D.L., Sprunger L.K.,

RA Kohrman D.C., Weisler M.H.;

RT "Exon organization, coding sequence, physical mapping, and polymorphic

RT intragenic markers for the human neuronal sodium channel gene SCN8A.";

RL Genomics 54:287-296 (1998).

[2]

RP SEQUENCE FROM N.A. (ISOFORM 2), TISSUE SPECIFICITY, AND DISEASE.
 RC STRAIN=057BL/60; TISSUE=Brain;
 RX MEDLINE=95400328; PubMed=7670495;
 RA Burgess D.L., Kohman D.C., Galt J., Plummer N.W., Jones J.M.,
 RT "Mutation of a new sodium channel gene, Scn8a, in the mouse mutant
 RL 'motor endplate disease'.";
 RN Nat. Genet. 10:461-465(1995).
 [3]
 RP SEQUENCE OF 93-205 FROM N.A., AND DISEASE.
 RC STRAIN=129/Sv; TISSUE=Brain;
 RX MEDLINE=96291923; PubMed=8663325;
 RA Kohman D.C., Harris J.B., Weisler M.H.,
 RT "Mutation detection in the med and medd alleles of the sodium channel
 RL Scn8a. Unusual splicing due to a minor class AT-AC intron.";
 RN J. Biol. Chem. 271:17576-17581(1996).
 [4]
 RP SEQUENCE OF 1411-1686 FROM N.A.
 RA Fan Z., Kyle J.W., Maklefski J.C.,
 RT "A putative novel Na channel alpha subunit cDNA isolated from mouse
 RL NB2a neuroblastoma cells.";
 RN Submitted (MAR-1995) to the EMBL/GenBank/DBJ databases.
 [5]
 RP ALTERNATIVE SPLICING (ISOFORMS 1; 4 AND 5).
 RC TISSUE=Brain, and fetal brain;
 RX MEDLINE=97442476; PubMed=9295353;
 RA Plummer N.W., McBurney M.W., Weisler M.H.,
 RT "Alternative splicing of the sodium channel SCN8a predicts a truncated
 RL two-domain protein in fetal brain and non-neuronal cells.";
 RN J. Biol. Chem. 272:24008-24015(1997).
 [6]
 RP VARIANT MEDJO THR-1317, AND VARIANT LEU-5.
 RC STRAIN=DBA/2MyD1;
 RX MEDLINE=96424513; PubMed=8815882;
 RA Kohman D.C., Smith M.R., Goldin A.L., Harris J., Weisler M.H.,
 RT "A missense mutation in the sodium channel Scn8a is responsible for
 RL cerebellar ataxia in the mouse mutant jolting.";
 RN J. Neurosci. 16:5993-5999(1996).
 [7]
 RP DISEASE.
 RX MEDLINE=21423786; PubMed=11532991;
 RA De Repentigny Y., Core P.D., Pool M., Bernier G., Girard S.,
 RT Vidal S.M., Kochary R.,
 RL "Pathological and genetic analysis of the degenerating muscle (dmu)
 Hum. Mol. Genet. 10:1819-1827(2001)."
 CC -1- FUNCTION: Mediates the voltage-dependent sodium ion permeability
 response to the voltage difference across the membrane, the
 protein forms a sodium-selective channel through which Na(+) ions
 may pass in accordance with their electrochemical gradient.
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
 CC -1- ALTERNATIVE PRODUCTS:
 CC Event=Alternative splicing; Named isoforms=5;
 CC Name=1; Synonyms=18a;
 CC IsoId=Q9WTU3-1; Sequence=Displayed;
 CC Name=2;
 CC IsoId=Q9WTU3-2; Sequence=VSP_050594;
 CC Name=3;
 CC IsoId=Q9WTU3-3; Sequence=VSP_050595;
 CC Name=4; Synonyms=18n;
 CC IsoId=Q9WTU3-4; Sequence=VSP_050596, VSP_050597;
 CC Name=5;
 CC IsoId=Q9WTU3-5; Sequence=VSP_050598;
 CC -1- TISSUE SPECIFICITY: Expressed in brain, cerebellum and spinal
 cord.
 CC -1- DOMAIN: The sequence contains 4 internal repeats, each with 5
 hydrophobic segments (S1,S2,S3,S5,S6) and one positively charged
 segment (S4). Segments S4 are probably the voltage-sensors and are
 characterized by a series of positively charged amino acids at
 every third position.
 CC -1- DISEASE: Defects in Scn8a are the cause of motor endplate disease
 (med). Med is a recessive neuromuscular disorder that is

CC characterized by lack of signal transmission at the neuromuscular
 CC junction, excess preterminal arborization and degeneration of
 CC cerebellar Purkinje cells. It produces early onset progressive
 CC paralysis of hind limbs, severe muscle atrophy and juvenile
 CC lethality.
 CC -1- DISEASE: Defects in Scn8a are the cause of the jolting mutant
 CC (medjo), a mild form of motor endplate disease which is
 CC characterized by the absence of spontaneous, regular, simple
 CC discharges from Purkinje cells. After 3 weeks of age, jolting mice
 CC are unsteady and have wide-based gait and a rhythmic tremor of
 CC head and neck induced by attempted movement.
 CC -1- DISEASE: Defects in Scn8a are a cause of degenerating muscle
 CC (dmu). Dmu is an autosomal recessive neuromuscular disorder that
 CC is characterized by skeletal and cardiac muscle degeneration. It
 CC produces early onset progressive loss of mobility of the hind
 CC limbs and subsequent lethality in the first month of life.
 CC -1- SIMILARITY: Belongs to the sodium channel family.
 CC -1- SIMILARITY: Contains 1 IQ domain.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
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 CC or_send_an_email_to_license@isb-sib.ch](http://www.isb-sib.ch/announce/or_send_an_email_to_license@isb-sib.ch)).
 CC -----
 CC EMBL, AF049617; AAD20438.1; -
 CC EMBL, U26707; AAC52242.1; -
 CC EMBL, U59964; AAC52708.1; -
 CC EMBL, U59963; AAC52708.1; JOINED.
 CC EMBL, U23158; AAA65599.1; -
 CC HSSP, P04775; 1BYX.
 CC MGD, MGI:103169; Scn8a.
 CC GO, GO:0001518; C:voltage-gated sodium channel complex; IC.
 CC GO, GO:0005248; F:voltage-gated sodium channel activity; MAS.
 CC GO, GO:0007628; P:adult walking behavior; IMP.
 CC GO, GO:0006814; P:sodium ion transport; NAS.
 CC InterPro: IPR001682; Ca/Na pore.
 CC InterPro: IPR002111; Cat_channel_TrpL.
 CC InterPro: IPR005821; Ion_trans.
 CC InterPro: IPR000048; IQ_region.
 CC InterPro: IPR005820; M-channel_nlg.
 CC InterPro: IPR001696; Na_channel.
 CC InterPro: IPR008054; Na_channel18.
 CC InterPro: IPR010526; Na_trans_assoc.
 CC Pfam, PF00520; Ion_trans; 4.
 CC Pfam, PF00612; IQ_1.
 CC Pfam, PF06512; Na_trans_assoc; 1.
 CC PRINTS, PR00170; NACHANNEL.
 CC PRINTS, PR01667; NACHANNEL8.
 CC PROSITE, PS00096; IQ; 1.
 CC Alternative splicing; ATP-binding; Disease mutation; Glycoprotein;
 CC Ion transport; Ionic channel; Multigene family; Polynorphism; Repeat;
 CC Sodium channel; Transmembrane; Voltage-gated channel.
 CC KW REPEAT 114 442 I.
 CC REPEAT 733 1005 II.
 CC REPEAT 1178 1493 III.
 CC REPEAT 1502 1799 IV.
 CC TRANSMEM 128 151 I.
 CC TRANSMEM 160 179 II.
 CC TRANSMEM 193 211 III.
 CC TRANSMEM 218 237 IV.
 CC TRANSMEM 253 277 V.
 CC TRANSMEM 388 413 VI.
 CC TRANSMEM 746 770 VII.
 CC TRANSMEM 782 805 VIII.
 CC TRANSMEM 814 833 IX.
 CC TRANSMEM 840 860 X.
 CC TRANSMEM 876 896 XI.
 CC TRANSMEM 950 975 XII.
 CC TRANSMEM 1192 1215 XIII.
 CC TRANSMEM 1229 1254 XIV.
 CC S2 of repeat I.
 CC S2 of repeat I.
 CC S3 of repeat I.
 CC S4 of repeat I.
 CC S5 of repeat I.
 CC S6 of repeat I.
 CC S1 of repeat II.
 CC S2 of repeat II.
 CC S3 of repeat II.
 CC S4 of repeat II.
 CC S5 of repeat II.
 CC S6 of repeat II.
 CC S1 of repeat III.
 CC S2 of repeat III.

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FT TRANSMEM 1261 1282 S3 of repeat III.
FT TRANSMEM 1267 1308 S4 of repeat III.
FT TRANSMEM 1328 1349 S5 of repeat III.
FT TRANSMEM 1436 1462 S6 of repeat III.
FT TRANSMEM 1516 1539 S1 of repeat IV.
FT TRANSMEM 1551 1574 S2 of repeat IV.
FT TRANSMEM 1581 1604 S3 of repeat IV.
FT TRANSMEM 1615 1636 S4 of repeat IV.
FT TRANSMEM 1652 1674 S5 of repeat IV.
FT TRANSMEM 1740 1764 S6 of repeat IV.
FT DOMAIN 1893 1932 IO.
FT NP_BIND 891 898 ATP (potential).
FT CARBOHYD 215 215 N-linked (GlcNAc. . .) (Potential).
FT CARBOHYD 289 289 N-linked (GlcNAc. . .) (Potential).
FT CARBOHYD 295 295 N-linked (GlcNAc. . .) (Potential).
FT CARBOHYD 308 308 N-linked (GlcNAc. . .) (Potential).
FT CARBOHYD 326 326 N-linked (GlcNAc. . .) (Potential).
FT CARBOHYD 344 344 N-linked (GlcNAc. . .) (Potential).
FT CARBOHYD 640 640 N-linked (GlcNAc. . .) (Potential).
FT CARBOHYD 875 875 N-linked (GlcNAc. . .) (Potential).
FT CARBOHYD 1045 1045 N-linked (GlcNAc. . .) (Potential).
FT CARBOHYD 1062 1062 N-linked (GlcNAc. . .) (Potential).
FT CARBOHYD 1089 1089 N-linked (GlcNAc. . .) (Potential).
FT CARBOHYD 1356 1356 N-linked (GlcNAc. . .) (Potential).
FT CARBOHYD 1370 1370 N-linked (GlcNAc. . .) (Potential).
FT CARBOHYD 1381 1381 N-linked (GlcNAc. . .) (Potential).
FT CARBOHYD 1766 1766 N-linked (GlcNAc. . .) (Potential).
FT VARSPLIC 428 673 Missing (in isoform 2).
FT VARSPLIC 664 664 /FTID=VSP_050594.
FT VARSPLIC 664 664 E -> EVKIDKATDS (in isoform 3).
FT VARSPLIC 1273 1280 /FTID=VSP_050595.
FT VARSPLIC 1273 1280 SLVSLIAN -> PLSISGLI (in isoform 4).

```

Query Match Best Local Similarity 89.3%; Score 25; DB 1; Length 1978; 100.0%; Pred. No. 1.5e-14; Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 4 VFMMVWVIGNLVVNLFLALLSSSF 28
Db 954 VFMMVWVIGNLVVNLFLALLSSSF 978

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RESULT 17
O88420 PRELIMINARY; PRT; 1978 AA.
AC O88420:
DT 01-NOV-1998 (TREMBLrel. 08, Created)
DT 01-NOV-1998 (TREMBLrel. 08, Last sequence update)
DT 01-MAR-2004 (TREMBLrel. 26, Last annotation update)
DE Voltage-gated sodium channel rRNA.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Dorsal root ganglia;
RX MEDLINE=96264329; PubMed=9603190;
RA Dietrich P.S., McGivern J.G., Delgado S.G., Koch B.D., Eglen R.M.,
RA Hunter J.C., Sangameswaran L.;
RT "Functional analysis of a voltage-gated sodium channel and its splice
RT variant from rat dorsal root ganglia.";
RL J. Neurochem. 70:2262-2272(1998).
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
CC -1- SIMILARITY: Belongs to the sodium channel family.
DR HSSP; P04775; IBBY.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0005151; C:voltage-gated sodium channel complex; IEA.
DR GO; GO:0005261; F:cation channel activity; IEA.
DR GO; GO:0005246; F:voltage-gated sodium channel activity; IEA.
DR GO; GO:0006812; P:cation transport; IEA.
DR GO; GO:0006814; P:sodium ion transport; IEA.

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DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR002111; Cat_channel_TrypL.
DR InterPro; IPR000183; Decarboxylase2.
DR InterPro; IPR005821; Ion trans.
DR InterPro; IPR000048; IQ_region.
DR InterPro; IPR005820; M+channel_n1g.
DR InterPro; IPR001696; Na_channel1.
DR InterPro; IPR008054; Na_channel8.
DR InterPro; IPR010526; Na_trans_assoc.
DR Pfam; PR00520; Ion_trans; 4.
DR Pfam; PR00612; IQ_1.
DR Pfam; PR06512; Na_trans_assoc; 1.
DR PRINTS; PR00170; NACHANNEL.
DR PRINTS; PR01667; NACHANNEL8.
DR SMART; SM00015; IQ; 1.
DR PROSITE; PS00096; IQ; 1.
DR PROSITE; PS00878; ODR_DC_2_1; UNKNOWN_1.
KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
KW Transport; Voltage-gated channel.
SQ SEQUENCE 1978 AA; 225157 MW; 9160843C5935B80B CRC64;

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Query Match Best Local Similarity 89.3%; Score 25; DB 2; Length 1978; 100.0%; Pred. No. 1.5e-14; Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 4 VFMMVWVIGNLVVNLFLALLSSSF 28
Db 954 VFMMVWVIGNLVVNLFLALLSSSF 978

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RESULT 18
CIN8 HUMAN STANDARD; PRT; 1980 AA.
ID CIN8_HUMAN
AC Q9UD0; Q95788; Q9NYX2; Q9UPB2;
DT 10-OCT-2003 (Rel. 42, Created)
DT 10-OCT-2003 (Rel. 42, Last sequence update)
DT 05-JUL-2004 (Rel. 44, Last annotation update)
DE Sodium channel protein type VIII alpha subunit (Voltage-gated sodium
DE channel alpha subunit Nav1.6).
GN Name=SCN8A;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A. (ISOFORM 4).
RC TISSUE=Brain, and Fetal brain;
RX MEDLINE=97442476; PubMed=9295353;
RA Plummer N.W., McBurney M.W., Weisler M.H.;
RT "Alternative splicing of the sodium channel SCN8A predicts a truncated
RT two-domain protein in fetal brain and non-neuronal cells.";
RL J. Biol. Chem. 272:24008-24015(1997).
RN [2]
RP SEQUENCE FROM N.A. (ISOFORMS 1; 2 AND 3).
RX MEDLINE=99047535; PubMed=9828131;
RA Plummer N.W., Galt J., Jones J.M., Burgess D.L., Sprunger L.K.,
RA Kohman D.C., Weisler M.H.;
RT "Exon organization, coding sequence, physical mapping, and polymorphic
RT intragenic markers for the human neuronal sodium channel gene SCN8A.";
RL Genomics 54:287-296(1998).
RN [3]
RP SEQUENCE FROM N.A. (ISOFORM 1).
RA Iin C., Numakura C., Kiyoshi H.;
RL Submitted (JUN-1999) to the EMBL/GenBank/DBJ databases.
RN [4]
RP SEQUENCE FROM N.A. (ISOFORM 1).
RA Jeong S.-Y., Goto J., Kanazawa I.;
RT "Cloning of cDNA for human voltage-gated sodium channel alpha subunit,
RT SCN8A.";
CC -1- FUNCTION: Mediates the EMBL/GenBank/DBJ databases.
CC of excitable membranes. Assuming opened or closed conformations in
CC response to the voltage difference across the membrane, the

```


FT CONFLICT 1445 1445 V -> I (in Ref. 1).
 FT CONFLICT 1519 1519 V -> I (in Ref. 1).
 FT CONFLICT 1702 1702 T -> A (in Ref. 4).
 SQ SEQUENCE 1980 AA; 225279 MW; 0BFC7BF137FD4F0 CRC64;

Query Match 89.3%; Score 25; DB 1; Length 1980;
 Best Local Similarity 100.0%; Pred. No. 1.5e-14;
 Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 4 VFMWVWVIGNLVNLFLALLSSSF 28
 Db 956 VFMWVWVIGNLVNLFLALLSSSF 980

RESULT 19

088421 PRELIMINARY; PRT; 1988 AA.
 ID 088421
 AC 088421
 DT 01-NOV-1998 (TREMBlrel. 08, Created)
 DT 01-NOV-1998 (TREMBlrel. 08, Last sequence update)
 DT 01-MAR-2004 (TREMBlrel. 26, Last annotation update)
 DE Voltage-gated sodium channel variant rRNA.
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 OC NCBI_TaxId=10116;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Dorsal root ganglia;
 RX MEDLINE=98264329; PubMed=9603190;
 RA Dierlich J.P., McGovern J.G., Delgado S.G., Koch B.D., Eglén R.M.,
 RA Hunter J.C., Sangameswaran L.;
 RT "Functional analysis of a voltage-gated sodium channel and its splice
 variant from rat dorsal root ganglia.";
 RL J. Neurochem. 70:2262-2272(1998).
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
 CC -1- SIMILARITY: Belongs to the sodium channel family.
 DR EMBL: AF049240; AAC26015.1; -.
 DR HSRP: P04775; IBYX.
 DR GO: GO:0016021; C:Integral to membrane; IEA.
 DR GO: GO:001518; C:Voltage-gated sodium channel complex; IEA.
 DR GO: GO:0005248; F:Voltage-gated sodium channel activity; IEA.
 DR GO: GO:0005248; F:Voltage-gated sodium channel activity; IEA.
 DR GO: GO:0006814; P:sodium ion transport; IEA.
 DR GO: GO:0006814; P:sodium ion transport; IEA.
 DR InterPro: IPR001682; Ca/Na_pore.
 DR InterPro: IPR002111; Cat_channel_TrtL.
 DR InterPro: IPR005821; Decarboxylase2.
 DR InterPro: IPR005821; Ion_trans.
 DR InterPro: IPR005820; M_channel_nlg.
 DR InterPro: IPR001696; Na_channel.
 DR InterPro: IPR008054; Na_channel18.
 DR InterPro: IPR010526; Na_trans_assoc.
 DR Pfam: PF00520; Ion_trans_4.
 DR Pfam: PF00612; IQ_1.
 DR Pfam: PF06512; Na_trans_assoc; 1.
 DR PRINTS: PRO0170; NACHANNEL.
 DR PRINTS: PRO1667; NACHANNEL8.
 DR SMART: SM00015; IQ_1.
 DR PROSITE: PS00878; ODR_DC_2_1; UNKNOWN 1.
 KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
 KW Transport; Voltage-gated channel.
 SQ SEQUENCE 1988 AA; 226186 MW; C1F7D87FDC266C8F CRC64;

Query Match 89.3%; Score 25; DB 2; Length 1988;
 Best Local Similarity 100.0%; Pred. No. 1.6e-14;
 Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 4 VFMWVWVIGNLVNLFLALLSSSF 28
 Db 964 VFMWVWVIGNLVNLFLALLSSSF 988

RESULT 20

015858 PRELIMINARY; PRT; 1977 AA.
 ID 015858
 AC 015858
 DT 01-NOV-1996 (TREMBlrel. 01, Created)
 DT 01-NOV-1996 (TREMBlrel. 01, Last sequence update)
 DT 01-MAR-2004 (TREMBlrel. 26, Last annotation update)
 DE Sodium channel alpha subunit.
 GN Name=hne-Na;
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OC NCBI_TaxId=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Thyroid;
 RX MEDLINE=95237189; PubMed=7720699;
 RA Klugbauer N., Lacinova L., Flocke V., Hofmann F.;
 RT "Structure and functional expression of a new member of the
 RT tetrodotoxin-sensitive voltage-activated sodium channel family from
 RT human neuroendocrine cells.";
 RL EMBO J. 14:1084-1090(1995).
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
 CC -1- SIMILARITY: Belongs to the sodium channel family.
 DR EMBL: X82835; CAA58042.1; -.
 DR PIR: S54771; S54771.
 DR HSRP: P04775; IBYX.
 DR GeneW: HGNC:10597; SCN9A.
 DR GO: GO:0005248; F:Voltage-gated sodium channel activity; TAS.
 DR GO: GO:0006814; P:sodium ion transport; TAS.
 DR InterPro: IPR001682; Ca/Na_pore.
 DR InterPro: IPR002111; Cat_channel_TrtL.
 DR InterPro: IPR005821; Ion_trans.
 DR InterPro: IPR000048; IQ_region.
 DR InterPro: IPR005820; M_channel_nlg.
 DR InterPro: IPR001696; Na_channel1.
 DR InterPro: IPR010526; Na_trans_assoc.
 DR Pfam: PF00520; Ion_trans_4.
 DR Pfam: PF00612; IQ_1.
 DR Pfam: PF06512; Na_trans_assoc; 1.
 DR PRINTS: PRO0170; NACHANNEL.
 DR SMART: SM00015; IQ_1.
 KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
 KW Transport; Voltage-gated channel.
 SQ SEQUENCE 1977 AA; 225195 MW; 17D67C8C32BC15FB CRC64;

Query Match 82.1%; Score 23; DB 2; Length 1977;
 Best Local Similarity 100.0%; Pred. No. 1.1e-12;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 6 MMWVWVIGNLVNLFLALLSSSF 28
 Db 938 MMWVWVIGNLVNLFLALLSSSF 960

RESULT 21

028644 PRELIMINARY; PRT; 1984 AA.
 ID 028644
 AC 028644
 DT 01-NOV-1996 (TREMBlrel. 01, Created)
 DT 01-NOV-1996 (TREMBlrel. 01, Last sequence update)
 DT 01-MAR-2004 (TREMBlrel. 26, Last annotation update)
 DE Sodium channel alpha-subunit.
 OS Oryctolagus cuniculus (Rabbit).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Lagomorpha; Leporidae; Oryctolagus.
 OC NCBI_TaxId=9986;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=New Zealand White; TISSUE=Sciatic nerve;
 RX MEDLINE=96074641; PubMed=7479311;

RA Belcher S.M., Zerillo C.A., Levenson R., Ritchie J.M., Howe J.R.;
 RT "Cloning of a sodium channel alpha subunit from rabbit Schwann
 cells.";
 RT Proc. Natl. Acad. Sci. U.S.A. 92:11034-11038(1995).
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
 CC -1- SIMILARITY: Belongs to the sodium channel family.
 DR EMBL: U55238; AA09159.1; -.
 DR HSSP: P04775; IBYI.
 DR GO: GO:0016021; C: integral to membrane; IEA.
 DR GO: GO:0001518; C: voltage-gated sodium channel complex; IEA.
 DR GO: GO:0005261; F: cation channel activity; IEA.
 DR GO: GO:0005248; F: voltage-gated sodium channel activity; IEA.
 DR GO: GO:0006812; P: cation transport; IEA.
 DR GO: GO:0006814; P: sodium ion transport; IEA.
 DR InterPro: IPR001682; Ca/Na_pore.
 DR InterPro: IPR002111; Cat_channel_TrpL.
 DR InterPro: IPR005821; Ion_trans.
 DR InterPro: IPR000948; IQ_region.
 DR InterPro: IPR005820; M_channel_nlg.
 DR InterPro: IPR001696; Na_channel.
 DR Pfam: PF00520; Ion_trans; 4.
 DR Pfam: PF06512; IQ; 1.
 DR Pfam: PF06512; Na_trans_assoc; 1.
 DR PRINTS: PR00170; NACHANNEL.
 DR SMART: SM00015; IQ; 1.
 KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
 KW Transport; Voltage-gated channel.
 SQ SEQUENCE 1984 AA; 225748 MW; 98F76860C9866AA0 CRC64;

Query Match 82.1%; Score 23; DB 2; Length 1984;
 Best Local Similarity 100.0%; Pred. No. 1.le-12;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 6 MMWVIGNLVYLNFLALILSSSF 28
 DB 946 MMWVIGNLVYLNFLALILSSSF 968

RESULT 22
 ID 008562; PRELIMINARY; PRT; 1984 AA.
 AC 008562;
 DT 01-JUN-1997 (TREMBLrel. 04, Created)
 DT 01-JUL-1997 (TREMBLrel. 04, Last sequence update)
 DT 05-JUL-2004 (TREMBLrel. 27, Last annotation update)
 DE PNI (Voltage-gated sodium channel) (Fragment).
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Rattus.
 OC NCBI_TaxID=10116;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA MEDLINE=97188502; PubMed=9037087;
 RA Toledo-Arai J.J., Moss B.L., He Z.J., Kozowski A.G., Whisenand T.,
 RA Levenson S.R., Wolf J.J., Silos-Santiago I., Halegoua S., Mandel G.;
 RA "Identification of PNI, a predominant voltage-dependent sodium channel
 RT expressed principally in peripheral neurons.";
 RT Proc. Natl. Acad. Sci. U.S.A. 94:1527-1532(1997).
 RL [2]
 RP SEQUENCE FROM N.A.
 RA MEDLINE=97007982; PubMed=8854872;
 RA Kozak C.A., Sangameswaran L.;
 RA "Genetic mapping of the peripheral sodium channel genes, Scn9a and
 RT Scn10a, in the mouse.";
 RL Mamm. Genome 7:787-788(1996).
 RN [3]
 RP SEQUENCE FROM N.A.
 RA Sangameswaran L., Fish L.M., Koch B.D., Rabert D.K., Delgado S.G.,
 RA Ilancka M., Jakeman L.B., Novakovic S., Wong K., Sze P., Tzoumaka E.,
 RA Stewart G.R., Herman R.C., Chan H., Eglyen R.M., Hunter J.C.;
 RA "A novel tetrodotoxin-sensitive, voltage-gated sodium channel
 RT expressed in rat and human dorsal root ganglia.";

RL J. Biol. Chem. 0:0-0(1997).
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
 CC -1- SIMILARITY: Belongs to the sodium channel family.
 DR EMBL: U79568; AA850403.1; -.
 DR EMBL: AF000368; AA880701.1; -.
 DR HSSP: P04775; IBYI.
 DR GO: GO:0016021; C: integral to membrane; IEA.
 DR GO: GO:0001518; C: voltage-gated sodium channel complex; IEA.
 DR GO: GO:0005261; F: cation channel activity; IEA.
 DR GO: GO:0005248; F: voltage-gated sodium channel activity; IEA.
 DR GO: GO:0006812; P: cation transport; IEA.
 DR GO: GO:0006814; P: sodium ion transport; IEA.
 DR InterPro: IPR001682; Ca/Na_pore.
 DR InterPro: IPR002111; Cat_channel_TrpL.
 DR InterPro: IPR005821; Ion_trans.
 DR InterPro: IPR000948; IQ_region.
 DR InterPro: IPR005820; M_channel_nlg.
 DR InterPro: IPR001696; Na_channel.
 DR Pfam: PF00520; Ion_trans; 4.
 DR Pfam: PF06512; IQ; 1.
 DR Pfam: PF06512; Na_trans_assoc; 1.
 DR PRINTS: PR00170; NACHANNEL.
 DR SMART: SM00015; IQ; 1.
 KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
 KW Transport; Voltage-gated channel.
 FT NON TER 1984 1984
 SQ SEQUENCE 1984 AA; 226037 MW; 386C38B9B5097091 CRC64;

Query Match 82.1%; Score 23; DB 2; Length 1984;
 Best Local Similarity 100.0%; Pred. No. 1.le-12;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 6 MMWVIGNLVYLNFLALILSSSF 28
 DB 948 MMWVIGNLVYLNFLALILSSSF 970

RESULT 23
 ID 080079; PRELIMINARY; PRT; 473 AA.
 AC 080079;
 DT 01-JUN-2003 (TREMBLrel. 24, Created)
 DT 01-JUN-2003 (TREMBLrel. 24, Last sequence update)
 DT 01-MAR-2004 (TREMBLrel. 26, Last annotation update)
 DE Sodium channel 6 (Fragment).
 OS Ictalurus punctatus (Channel catfish).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Siluriformes;
 OC Ictaluridae; Ictalurus.
 OC NCBI_TaxID=7998;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Lu Y., Lopreato G.F., Zakon H.H.;
 RA Submitted (DEC-2002) to the EMBL/GenBank/DBJ databases.
 RA EMBL: AY204537; AAC060425.1; -.
 DR GO: GO:0016021; C: integral to membrane; IEA.
 DR GO: GO:0005261; F: cation channel activity; IEA.
 DR GO: GO:0006812; P: cation transport; IEA.
 DR GO: GO:0006814; P: sodium ion transport; IEA.
 DR InterPro: IPR001682; Ca/Na_pore.
 DR InterPro: IPR005821; Ion_trans.
 DR InterPro: IPR005820; M_channel_nlg.
 DR InterPro: IPR001696; Na_channel.
 DR Pfam: PF00520; Ion_trans; 1.
 DR Pfam: PF06512; Na_trans_assoc; 1.
 KW Ion transport; Ionic channel; Transmembrane; Transport.
 FT NON TER 473 473
 FT NON TER 1 1
 SQ SEQUENCE 473 AA; 53617 MW; D922504FD1CODF1B CRC64;
 Query Match 78.6%; Score 22; DB 2; Length 473;
 Best Local Similarity 100.0%; Pred. No. 3e-12;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 7 MVMVIGNLVNLFLLALISF 28
 |||||
 DB 12 MVMVIGNLVNLFLLALISF 33

RESULT 24

Q800U1 PRELIMINARY; PRT; 488 AA.

ID Q800U1
 AC Q800U1;
 DT 01-JUN-2003 (TEMBLrel. 24, Created)
 DT 01-JUN-2003 (TEMBLrel. 24, last sequence update)
 DT 01-MAR-2004 (TEMBLrel. 26, last annotation update)
 DE Sodium channel 4 (Fragment).
 OS Ictalurus punctatus (Channel catfish).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Siluriformes;
 OC Ictaluridae; Ictalurus.
 OC NCBI_TaxId=7998;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Lu Y., Lopreato G.F., Zakon H.H.;
 RL Submitted (DEC-2002) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AY204535; AA060423.1; -
 DR GO; GO:0016021; C:integral to membrane; IEA.
 DR GO; GO:0005261; P:cation channel activity; IEA.
 DR GO; GO:0006812; P:cation transport; IEA.
 DR InterPro; IPR001682; C8/Na_pore.
 DR InterPro; IPR005821; Ion_trans.
 DR InterPro; IPR005820; M+channel_nlg.
 DR InterPro; IPR010526; Na_trans_assoc.
 DR Pfam; PF00520; Ion_trans. 1.
 DR Pfam; PF06512; Na_trans_assoc. 1.
 DR Ion transport; Ionic channel; Transmembrane; Transport.
 KM NON_TER 1
 FT NON_TER 1
 SQ SEQUENCE 488 AA; 55100 MW; 379A1BAA105F3A8F CRC64;

Query Match 78.6%; Score 22; DB 2; Length 488;
 Best Local Similarity 100.0%; Pred. No. 3.1e-12;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 7 MVMVIGNLVNLFLLALISF 28
 |||||
 DB 12 MVMVIGNLVNLFLLALISF 33

RESULT 25

Q90227 PRELIMINARY; PRT; 744 AA.

ID Q90227
 AC Q90227;
 DT 01-DEC-2001 (TEMBLrel. 19, Created)
 DT 01-DEC-2001 (TEMBLrel. 19, last sequence update)
 DT 01-MAR-2004 (TEMBLrel. 26, last annotation update)
 DE Sodium channel 4 (Fragment).
 OS Sternopygus macrurus.
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Gymnotiformes;
 OC Sternopygidae; Sternopygus.
 OC NCBI_TaxId=77841;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA MEDLINE=21310016; PubMed=11416226;
 RA Lopreato G.F., Lu Y., Southwell A., Atkinson N.S., Hillis D.M.,
 RA Wilcox T.P., Zakon H.H.;
 RT "Evolution and divergence of sodium channel genes in vertebrates.";
 RL Proc. Natl. Acad. Sci. U.S.A. 98:7588-7592(2001).
 DR EMBL; AF378142; AKS5440.1; -
 DR GO; GO:0016021; C:integral to membrane; IEA.
 DR GO; GO:0005261; P:cation channel activity; IEA.
 DR GO; GO:0006812; P:cation transport; IEA.
 DR InterPro; IPR001682; C8/Na_pore.
 DR InterPro; IPR002111; Cat_channel_TrpL.

DR InterPro; IPR005821; Ion_trans.
 DR InterPro; IPR005820; M+channel_nlg.
 DR InterPro; IPR010526; Na_trans_assoc.
 DR Pfam; PF00520; Ion_trans. 2.
 DR Pfam; PF06512; Na_trans_assoc. 1.
 DR Ion transport; Ionic channel; Transmembrane; Transport.
 KM NON_TER 1
 FT NON_TER 1
 SQ SEQUENCE 744 AA; 84557 MW; B0D6E7B203893C28 CRC64;

Query Match 78.6%; Score 22; DB 2; Length 744;
 Best Local Similarity 100.0%; Pred. No. 4.4e-12;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 7 MVMVIGNLVNLFLLALISF 28
 |||||
 DB 184 MVMVIGNLVNLFLLALISF 205

Search completed: January 27, 2005, 17:51:32
 Job time : 94.5 secs

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OM protein - protein search, using sw model

Run on: January 27, 2005, 17:36:50 ; Search time 22.5 Seconds
(without alignments)
82.529 Million cell updates/sec

Title: US-10-608-584-13
Perfect score: 28
Sequence: 1 CLTFVMMVWVIGNLVNLFLALLSSF 28

Scoring table:
Gapop 60.0 , Gapext 60.0

Searched: 478139 seqs, 66318000 residues

Word size : 0

Total number of hits satisfying chosen parameters: 478139

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Listing first 100 summaries

Database :
1: Issued Patents AA:*
2: /cgn2_6/ptodata/1/1aa/5A_COMB.pep:*
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4: /cgn2_6/ptodata/1/1aa/6A_COMB.pep:*
5: /cgn2_6/ptodata/1/1aa/6B_COMB.pep:*
6: /cgn2_6/ptodata/1/1aa/6C_COMB.pep:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
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2	28	100.0	2005	4	US-09-457-571-7
3	25	89.3	1976	3	US-09-024-020B-9
4	25	89.3	1976	3	US-09-425-043-9
5	25	89.3	1978	3	US-09-024-020B-3
6	25	89.3	1978	3	US-09-425-043-3
7	25	89.3	1988	3	US-09-024-020B-4
8	25	89.3	1988	3	US-09-425-043-4
9	23	82.1	1835	3	US-08-836-325-15
10	23	82.1	1835	3	US-09-457-571-15
11	23	82.1	1969	4	US-08-836-325-16
12	23	82.1	1969	4	US-09-457-571-16
13	23	82.1	1977	4	US-09-976-594-757
14	23	82.1	1977	4	US-09-919-039-367
15	23	82.1	1984	4	US-08-836-325-10
16	23	82.1	1984	4	US-09-457-571-10
17	23	82.1	1989	3	US-08-836-325-11
18	23	82.1	1989	3	US-08-836-325-12
19	23	82.1	1989	4	US-09-457-571-11
20	23	82.1	1989	4	US-09-457-571-12
21	22	78.6	1836	4	US-10-162-012-24
22	22	75.0	1024	4	US-09-562-737-81
23	21	75.0	2016	4	US-09-634-920-4
24	21	75.0	2016	4	US-09-514-907A-2
25	21	75.0	2016	4	US-09-896-994-2
26	21	75.0	2016	4	US-09-840-125-4
27	17	60.7	1820	3	US-07-998-289B-8

28	17	60.7	2100	2	US-08-808-793-23	Sequence 23, Appl
29	17	60.7	2100	3	US-08-772-512A-19	Sequence 19, Appl
30	17	60.7	2105	3	US-08-808-793-3	Sequence 3, Appl
31	17	60.7	2105	3	US-08-772-512A-3	Sequence 3, Appl
32	16	57.1	813	4	US-08-836-325-8	Sequence 8, Appl
33	16	57.1	813	4	US-09-457-571-8	Sequence 8, Appl
34	16	57.1	1011	3	US-08-836-325-2	Sequence 2, Appl
35	16	57.1	1011	4	US-09-457-571-2	Sequence 2, Appl
36	12	42.9	2104	3	US-08-808-793-4	Sequence 4, Appl
37	12	42.9	2104	3	US-08-772-512A-4	Sequence 4, Appl
38	9	32.1	1024	4	US-09-562-737-82	Sequence 82, Appl
39	9	32.1	1024	4	US-09-562-737-83	Sequence 83, Appl
40	9	32.1	1024	4	US-09-562-737-84	Sequence 84, Appl
41	9	32.1	1024	4	US-09-562-737-85	Sequence 85, Appl
42	9	32.1	1024	4	US-09-562-737-86	Sequence 86, Appl
43	9	32.1	1024	4	US-09-562-737-87	Sequence 87, Appl
44	9	32.1	1024	4	US-09-562-737-88	Sequence 88, Appl
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46	9	32.1	1024	4	US-09-562-737-90	Sequence 90, Appl
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48	9	32.1	1956	3	US-08-843-417-10	Sequence 10, Appl
49	9	32.1	1956	4	US-09-527-013-2	Sequence 2, Appl
50	9	32.1	1956	4	US-09-527-013-10	Sequence 10, Appl
51	9	32.1	1957	4	US-08-669-656A-2	Sequence 2, Appl
52	9	32.1	1957	4	US-08-669-656A-8	Sequence 8, Appl
53	9	32.1	2132	4	US-08-669-656A-6	Sequence 6, Appl
54	8	28.6	501	1	US-08-331-394-4	Sequence 4, Appl
55	8	28.6	501	1	US-08-250-858-4	Sequence 4, Appl
56	8	28.6	501	1	US-08-446-915-4	Sequence 4, Appl
57	8	28.6	501	2	US-08-744-119-4	Sequence 4, Appl
58	8	28.6	501	4	US-08-779-599-4	Sequence 4, Appl
59	8	28.6	501	5	PCT-US95-06639-4	Sequence 4, Appl
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61	7	25.0	197	3	US-08-833-488B-4	Sequence 4, Appl
62	7	25.0	197	3	US-08-833-488B-9	Sequence 9, Appl
63	7	25.0	197	3	US-08-833-488B-28	Sequence 28, Appl
64	7	25.0	199	3	US-08-833-488B-14	Sequence 14, Appl
65	7	25.0	204	6	5187075-6	Patent No. 5187075
66	7	25.0	253	3	US-08-833-488B-20	Sequence 20, Appl
67	7	25.0	431	4	US-09-592-998C-9	Sequence 9, Appl
68	7	25.0	431	4	US-09-592-998C-10	Sequence 10, Appl
69	7	25.0	435	4	US-09-446-861-127	Sequence 127, App
70	7	25.0	461	3	US-09-122-210-2	Sequence 2, Appl
71	7	25.0	461	3	US-09-443-681-2	Sequence 2, Appl
72	7	25.0	573	4	US-09-643-657-18	Sequence 18, Appl
73	7	25.0	586	4	US-10-140-002-46	Sequence 46, Appl
74	7	25.0	591	4	US-09-643-657-3	Sequence 3, Appl
75	7	25.0	633	2	US-08-736-770-3	Sequence 3, Appl
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77	7	25.0	1023	4	US-10-140-002-200	Sequence 200, App
78	7	25.0	1233	4	US-09-354-147C-7	Sequence 7, Appl
79	7	25.0	1233	4	US-09-354-147C-8	Sequence 8, Appl
80	7	25.0	1498	2	US-08-404-531B-28	Sequence 28, Appl
81	7	25.0	1498	2	US-08-404-531B-29	Sequence 29, Appl
82	7	25.0	1498	3	US-08-476-900A-28	Sequence 28, Appl
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84	7	25.0	1498	3	US-08-488-546A-28	Sequence 28, Appl
85	7	25.0	1498	3	US-08-488-546A-29	Sequence 29, Appl
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87	7	25.0	1580	3	US-09-208-716-1	Sequence 1, Appl
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89	7	25.0	1581	3	US-08-476-900A-6	Sequence 6, Appl
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92	7	25.0	1581	3	US-08-726-320-4	Sequence 4, Appl
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94	7	25.0	1581	3	US-09-208-716-4	Sequence 4, Appl
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98	7	25.0	1582	3	US-08-726-320-5	Sequence 5, Appl
99	7	25.0	1582	3	US-09-208-716-5	Sequence 5, Appl
100	7	25.0	1765	4	US-09-354-147C-2	Sequence 2, Appl

ALIGNMENTS

RESULT 1

US-08-836-325-7
Sequence 7, Application US/08836325
Patent No. 6110672
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Halegoua, Simon
APPLICANT: Borden, Laurence A.
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
TITLE OF INVENTION: Thereof
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/836,325
FILING DATE: 2-MAY-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917.0240002
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2540
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 2005 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-08-836-325-7

Query Match 100.0%; Score 28; DB 3; Length 2005;
Best Local Similarity 100.0%; Pred. No. 1.1e-18;

Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CLTVFMWVWVIGNLVYVNLFLALLSSSF 28

Db 959 CLTVFMWVWVIGNLVYVNLFLALLSSSF 986

RESULT 2

US-09-457-571-7
Sequence 7, Application US/09457571
Patent No. 6703486

GENERAL INFORMATION:

APPLICANT: Mandel, Gail
APPLICANT: Halegoua, Simon
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
TITLE OF INVENTION: Thereof
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/457,571
FILING DATE: 09-DEC-1999
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/836,325
FILING DATE: 02-MAY-1997
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917.0240003
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 2005 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-457-571-7

Query Match 100.0%; Score 28; DB 4; Length 2005;
Best Local Similarity 100.0%; Pred. No. 1.1e-18;

Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CLTVFMWVWVIGNLVYVNLFLALLSSSF 28

Db 959 CLTVFMWVWVIGNLVYVNLFLALLSSSF 986

RESULT 3

US-09-024-020B-9
Sequence 9, Application US/09024020B
Patent No. 6030810

GENERAL INFORMATION:
APPLICANT: DELGADO, STEPHEN G.
APPLICANT: DIETRICH, PAUL S.
APPLICANT: FISH, LINDA M.
APPLICANT: HERMAN, RONALD C.
APPLICANT: SANGAMESWARAN, LAKSHMI

TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE

TITLE OF INVENTION: SODIUM CHANNEL I-SUBUNIT AND A SPLICE VARIANT THEREOF
NUMBER OF SEQUENCES: 43
CORRESPONDENCE ADDRESS:
ADDRESSEE: JANET PAULINE CLARK
STREET: 3401 HILLVIEW AVENUE, MS A2-250
CITY: PALO ALTO
STATE: CA
COUNTRY: U.S.A.
ZIP: 94304-1397
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/024,020B
FILING DATE: 16-FEB-1998
CLASSIFICATION: 536
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/039,447
FILING DATE: 26-FEB-1997
ATTORNEY/AGENT INFORMATION:
NAME: CLARK, JANET P.
REGISTRATION NUMBER: 34,799
REFERENCE/DOCKET NUMBER: R0020B-REG
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 852-3097
TELEFAX: (650) 855-5322
INFORMATION FOR SEQ ID NO: 9:
SEQUENCE CHARACTERISTICS:
LENGTH: 1976 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-024-020B-9

Query Match 89.3%; Score 25; DB 3; Length 1976;
Best Local Similarity 100.0%; Pred. No. 8.6e-16;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 VPMWVYIGNLVNLFLLALLSSF 28
Db 952 VPMWVYIGNLVNLFLLALLSSF 976

RESULT 4
US-09-425-043-9
Sequence 9, Application US/09425043
Patent No. 6335172
GENERAL INFORMATION:
APPLICANT: DELGADO, STEPHEN G.
APPLICANT: DIETRICH, PAUL S.
APPLICANT: FISH, LINDA M.
APPLICANT: HERMAN, RONALD C.
APPLICANT: SANGAMESWARAN, LAKSHMI
TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE
SODIUM CHANNEL I-SUBUNIT AND A SPLICE VARIANT THEREOF
NUMBER OF SEQUENCES: 43
CORRESPONDENCE ADDRESS:
ADDRESSEE: JANET PAULINE CLARK
STREET: 3401 HILLVIEW AVENUE, MS A2-250
CITY: PALO ALTO
STATE: CA
COUNTRY: U.S.A.
ZIP: 94304-1397
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/425,043

FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 09/024,020
FILING DATE: 16-FEB-1998
APPLICATION NUMBER: US 60/039,447
FILING DATE: 26-FEB-1997
ATTORNEY/AGENT INFORMATION:
NAME: CLARK, JANET P.
REGISTRATION NUMBER: 34,799
REFERENCE/DOCKET NUMBER: R0020B-REG
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 852-3097
TELEFAX: (650) 855-5322
INFORMATION FOR SEQ ID NO: 9:
SEQUENCE CHARACTERISTICS:
LENGTH: 1976 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-425-043-9

Query Match 89.3%; Score 25; DB 3; Length 1976;
Best Local Similarity 100.0%; Pred. No. 8.6e-16;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 VPMWVYIGNLVNLFLLALLSSF 28
Db 952 VPMWVYIGNLVNLFLLALLSSF 976

RESULT 5
US-09-024-020B-3
Sequence 3, Application US/09024020B
Patent No. 6030810
GENERAL INFORMATION:
APPLICANT: DELGADO, STEPHEN G.
APPLICANT: DIETRICH, PAUL S.
APPLICANT: FISH, LINDA M.
APPLICANT: HERMAN, RONALD C.
APPLICANT: SANGAMESWARAN, LAKSHMI
TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE
SODIUM CHANNEL I-SUBUNIT AND A SPLICE VARIANT THEREOF
NUMBER OF SEQUENCES: 43
CORRESPONDENCE ADDRESS:
ADDRESSEE: JANET PAULINE CLARK
STREET: 3401 HILLVIEW AVENUE, MS A2-250
CITY: PALO ALTO
STATE: CA
COUNTRY: U.S.A.
ZIP: 94304-1397
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/024,020B
FILING DATE: 16-FEB-1998
CLASSIFICATION: 536
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/039,447
FILING DATE: 26-FEB-1997
ATTORNEY/AGENT INFORMATION:
NAME: CLARK, JANET P.
REGISTRATION NUMBER: 34,799
REFERENCE/DOCKET NUMBER: R0020B-REG
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 852-3097
TELEFAX: (650) 855-5322
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:

LENGTH: 1978 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-024-020B-3

Query Match 89.3%; Score 25; DB 3; Length 1978;
Best Local Similarity 100.0%; Pred. No. 8.6e-16;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 VFMMVWVIGNLVNLFLALLSSSF 28
DB 954 VFMMVWVIGNLVNLFLALLSSSF 978

RESULT 6
US-09-425-043-3
Sequence 3, Application US/09425043
Patent No. 6335172
GENERAL INFORMATION:
APPLICANT: DELGADO, STEPHEN G.
APPLICANT: DIETRICH, PAUL S.
APPLICANT: FISH, LINDA W.
APPLICANT: HERMAN, RONALD C.
APPLICANT: SANGAMESWARAN, LAKSHMI
TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE
TITLE OF INVENTION: SODIUM CHANNEL I-SUBUNIT AND A SPLICE VARIANT THEREOF
NUMBER OF SEQUENCES: 43
CORRESPONDENCE ADDRESS:
ADDRESSEE: JANET PAULINE CLARK
STREET: 3401 HILLVIEW AVENUE, MS A2-250
CITY: PALO ALTO
STATE: CA
COUNTRY: U.S.A.
ZIP: 94304-1397
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/425,043
FILING DATE:
CLASSIFICATION:
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: US 09/024,020
FILING DATE: 16-FEB-1998
APPLICATION NUMBER: US 60/039,447
FILING DATE: 26-FEB-1997
ATTORNEY/AGENT INFORMATION:
NAME: CLARK, JANET P.
REGISTRATION NUMBER: 34,799
REFERENCE/DOCKET NUMBER: R0020B-REG
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 852-3097
TELEFAX: (650) 855-5322
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 1978 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-425-043-3

Query Match 89.3%; Score 25; DB 3; Length 1978;
Best Local Similarity 100.0%; Pred. No. 8.6e-16;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 4 VFMMVWVIGNLVNLFLALLSSSF 28
DB 954 VFMMVWVIGNLVNLFLALLSSSF 978

RESULT 7
US-09-024-020B-4
Sequence 4, Application US/09024020B
Patent No. 6030810
GENERAL INFORMATION:
APPLICANT: DELGADO, STEPHEN G.
APPLICANT: DIETRICH, PAUL S.
APPLICANT: FISH, LINDA W.
APPLICANT: HERMAN, RONALD C.
APPLICANT: SANGAMESWARAN, LAKSHMI
TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE
TITLE OF INVENTION: SODIUM CHANNEL I-SUBUNIT AND A SPLICE VARIANT THEREOF
NUMBER OF SEQUENCES: 43
CORRESPONDENCE ADDRESS:
ADDRESSEE: JANET PAULINE CLARK
STREET: 3401 HILLVIEW AVENUE, MS A2-250
CITY: PALO ALTO
STATE: CA
COUNTRY: U.S.A.
ZIP: 94304-1397
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/024,020B
FILING DATE: 16-FEB-1998
CLASSIFICATION: 536
PRIORITY APPLICATION DATA:
APPLICATION NUMBER: US 60/039,447
FILING DATE: 26-FEB-1997
ATTORNEY/AGENT INFORMATION:
NAME: CLARK, JANET P.
REGISTRATION NUMBER: 34,799
REFERENCE/DOCKET NUMBER: R0020B-REG
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 852-3097
TELEFAX: (650) 855-5322
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 1988 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-024-020B-4

Query Match 89.3%; Score 25; DB 3; Length 1988;
Best Local Similarity 100.0%; Pred. No. 8.6e-16;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 VFMMVWVIGNLVNLFLALLSSSF 28
DB 964 VFMMVWVIGNLVNLFLALLSSSF 988

RESULT 8
US-09-425-043-4
Sequence 4, Application US/09425043
Patent No. 6335172
GENERAL INFORMATION:
APPLICANT: DELGADO, STEPHEN G.
APPLICANT: DIETRICH, PAUL S.
APPLICANT: FISH, LINDA W.
APPLICANT: HERMAN, RONALD C.
APPLICANT: SANGAMESWARAN, LAKSHMI
TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE
TITLE OF INVENTION: SODIUM CHANNEL I-SUBUNIT AND A SPLICE VARIANT THEREOF
NUMBER OF SEQUENCES: 43
CORRESPONDENCE ADDRESS:


```

ADDRESS: JANET PAULINE CLARK
STREET: 3401 HILLVIEW AVENUE, MS A2-250
CITY: PALO ALTO
STATE: CA
COUNTRY: U.S.A.
ZIP: 94304-1397
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/425,043
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 09/024,020
FILING DATE: 16-FEB-1998
APPLICATION NUMBER: US 60/039,447
FILING DATE: 26-FEB-1997
ATTORNEY/AGENT INFORMATION:
NAME: CLARK, JANET P.
REGISTRATION NUMBER: 34,799
REFERENCE/DOCKET NUMBER: R0020B-REG
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 852-3097
TELEFAX: (650) 852-5322
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 1988 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-425-043-4

Query Match      89.3%; Score 25; DB 3; Length 1988;
Best Local Similarity 100.0%; Pred. No. 8,66-16;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 VFMVMVIGNLVNLFLALLISSF 28
DB 964 VFMVMVIGNLVNLFLALLISSF 988

RESULT 9
US-08-836-325-15
Sequence 15, Application US/08836325
Patent No. 6110672
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Halegoua, Simon
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Thereof, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational,
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
TITLE OF INVENTION: Thereof
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESS: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/836,325

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FILING DATE: 2-MAY-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917,0240002
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 15:
SEQUENCE CHARACTERISTICS:
LENGTH: 1835 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: not relevant
MOLECULE TYPE: protein
US-08-836-325-15

Query Match      82.1%; Score 23; DB 3; Length 1835;
Best Local Similarity 100.0%; Pred. No. 6,7e-14;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 MMVMVIGNLVNLFLALLISSF 28
DB 882 MMVMVIGNLVNLFLALLISSF 904

RESULT 10
US-09-457-571-15
Sequence 15, Application US/09457571
Patent No. 6703486
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Halegoua, Simon
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Thereof, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational,
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
TITLE OF INVENTION: Thereof
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESS: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/457,571
FILING DATE: 09-DEC-1999
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/836,325
FILING DATE: 02-MAY-1997
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401

```

FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917,0240003
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 15:
SEQUENCE CHARACTERISTICS:
LENGTH: 1835 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: not relevant
MOLECULE TYPE: protein
US-09-457-571-15

Query Match 82.1%; Score 23; DB 4; Length 1835;
Best Local Similarity 100.0%; Pred. No. 6,7e-14;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 6 MMVVVIGNLVVNLFLALLSSF 28
DB 882 MMVVVIGNLVVNLFLALLSSF 904

RESULT 11
US-08-836-325-16
Sequence 16, Application US/08836325
Patent No. 6110672
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Halegoua, Simon
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
TITLE OF INVENTION: Thereof
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/836,325
FILING DATE: 2-MAY-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917,0240002
TELECOMMUNICATION INFORMATION:

TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 16:
SEQUENCE CHARACTERISTICS:
LENGTH: 1969 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-836-325-16

Query Match 82.1%; Score 23; DB 3; Length 1969;
Best Local Similarity 100.0%; Pred. No. 7,2e-14;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 6 MMVVVIGNLVVNLFLALLSSF 28
DB 938 MMVVVIGNLVVNLFLALLSSF 960

RESULT 12
US-09-457-571-16
Sequence 16, Application US/09457571
Patent No. 6703486
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Halegoua, Simon
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
TITLE OF INVENTION: Thereof
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/457,571
FILING DATE: 09-DEC-1999
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/836,325
FILING DATE: 02-MAY-1997
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917,0240003
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 16:
SEQUENCE CHARACTERISTICS:
LENGTH: 1969 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant

TOPOLOGY: linear
MOLECULE TYPE: protein
US-09-457-571-16

Query Match 82.1%; Score 23; DB 4; Length 1969;
Best Local Similarity 100.0%; Pred. No. 7.2e-14;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 MMWVIGNLVVLNLFALLLSF 28
Db 938 MMWVIGNLVVLNLFALLLSF 960

RESULT 13
US-09-976-594-757
Sequence 757, Application US/09976594
Patent No. 6673549
GENERAL INFORMATION:
APPLICANT: Furness, Michael
APPLICANT: Buchbinder, Jenny
TITLE OF INVENTION: GENES EXPRESSED IN C3A LAYER CELL CULTURES TREATED WITH STEROIDS
FILE REFERENCE: PA-0041 US
CURRENT APPLICATION NUMBER: US/09/976,594
PRIOR FILING DATE: 2001-10-12
PRIOR APPLICATION NUMBER: 60/240,409
NUMBER OF SEQ ID NOS: 1143
SOFTWARE: PERL Program
SEQ ID NO 757
LENGTH: 1977
TYPE: PRT
ORGANISM: Homo sapiens
NAME/KEY: misc feature
OTHER INFORMATION: Incyte ID No. 6673549 1719478CD1
US-09-976-594-757

Query Match 82.1%; Score 23; DB 4; Length 1977;
Best Local Similarity 100.0%; Pred. No. 7.2e-14;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 MMWVIGNLVVLNLFALLLSF 28
Db 938 MMWVIGNLVVLNLFALLLSF 960

RESULT 14
US-09-919-039-367
Sequence 367, Application US/09919039
Patent No. 6727066
GENERAL INFORMATION:
APPLICANT: Kaeser, Matthew R.
TITLE OF INVENTION: GENES EXPRESSED IN TREATED HUMAN C3A LIVER CELL CULTURES
FILE REFERENCE: PA-0035 US
CURRENT APPLICATION NUMBER: US/09/919,039
PRIOR FILING DATE: 2002-09-09
PRIOR APPLICATION NUMBER: 60/222,113
NUMBER OF SEQ ID NOS: 401
SOFTWARE: PERL Program
SEQ ID NO 367
LENGTH: 1977
TYPE: PRT
ORGANISM: Homo sapiens
NAME/KEY: misc feature
OTHER INFORMATION: Incyte ID No. 6727066 1719478CD1
US-09-919-039-367

Query Match 82.1%; Score 23; DB 4; Length 1977;
Best Local Similarity 100.0%; Pred. No. 7.2e-14;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 MMWVIGNLVVLNLFALLLSF 28
Db 938 MMWVIGNLVVLNLFALLLSF 960

RESULT 15
US-08-836-325-10
Sequence 10, Application US/08836325
Patent No. 6110672
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Halegoua, Simon
APPLICANT: Borden, Laurence A.
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
TITLE OF INVENTION: Thereof
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC Compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/836,325
FILING DATE: 2-MAY-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917,0240002
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 10:
SEQUENCE CHARACTERISTICS:
LENGTH: 1984 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-836-325-10

Query Match 82.1%; Score 23; DB 3; Length 1984;
Best Local Similarity 100.0%; Pred. No. 7.2e-14;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 MMWVIGNLVVLNLFALLLSF 28
Db 948 MMWVIGNLVVLNLFALLLSF 970

RESULT 16
US-09-457-571-10
Sequence 10, Application US/09457571
Patent No. 6703486
GENERAL INFORMATION:

APPLICANT: Mandel, Gail
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
TITLE OF INVENTION: Thereof
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/457,571
FILING DATE: 09-DEC-1999
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/836,325
FILING DATE: 02-MAY-1997
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917.0240003
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 10:
SEQUENCE CHARACTERISTICS:
LENGTH: 1984 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-09-457-571-10

Query Match 82.1%; Score 23; DB 4; Length 1984;
Best Local Similarity 100.0%; Pred. No. 7.2e-14;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 6 MMWVIGNLVVNLFLALLISF 28
Db 948 MMWVIGNLVVNLFLALLISF 970

RESULT 17
US-08-836-325-11
Sequence 11, Application US/08836325
Patent No. 6110672
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Haleboua, Simon
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
TITLE OF INVENTION: Thereof

NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/836,325
FILING DATE: 2-MAY-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917.0240002
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 11:
SEQUENCE CHARACTERISTICS:
LENGTH: 1989 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: not relevant
MOLECULE TYPE: protein
US-08-836-325-11

Query Match 82.1%; Score 23; DB 3; Length 1989;
Best Local Similarity 100.0%; Pred. No. 7.2e-14;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 6 MMWVIGNLVVNLFLALLISF 28
Db 949 MMWVIGNLVVNLFLALLISF 971

RESULT 18
US-08-836-325-12
Sequence 12, Application US/08836325
Patent No. 6110672
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Haleboua, Simon
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
TITLE OF INVENTION: Thereof
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:

MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/836,325
FILING DATE: 2-MAY-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917,0240002
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 12:
SEQUENCE CHARACTERISTICS:
LENGTH: 1989 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: not relevant
MOLECULE TYPE: protein
US-08-836-325-12

Query March 82.1%; Score 23; DB 3; Length 1989;
Best Local Similarity 100.0%; Pred. No. 7.2e-14;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 MMWVIGNLVYLNFLALLLSF 28
DB 949 MMWVIGNLVYLNFLALLLSF 971

RESULT 19
US-09-457-571-11
Sequence 11, Application US/09457571
Patent No. 6703486
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Thereof, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
Thereof
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESSES:
ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N.W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/457,571
FILING DATE: 09-DEC-1999
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/836,325

FILING DATE: 02-MAY-1997
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917,0240003
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 11:
SEQUENCE CHARACTERISTICS:
LENGTH: 1989 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: not relevant
MOLECULE TYPE: protein
US-09-457-571-11

Query March 82.1%; Score 23; DB 4; Length 1989;
Best Local Similarity 100.0%; Pred. No. 7.2e-14;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 MMWVIGNLVYLNFLALLLSF 28
DB 949 MMWVIGNLVYLNFLALLLSF 971

RESULT 20
US-09-457-571-12
Sequence 12, Application US/09457571
Patent No. 6703486
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Thereof, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
Thereof
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESSES:
ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N.W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/457,571
FILING DATE: 09-DEC-1999
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/836,325
FILING DATE: 02-MAY-1997
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995

Fri Jan 28 09:32:00 2005

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PRIOR APPLICATION DATA: 08/334,029
APPLICATION NUMBER: 02-NOV-1994
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Iudwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917,0240003
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2540
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 12:
SEQUENCE CHARACTERISTICS:
LENGTH: 1989 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: not relevant
MOLECULE TYPE: protein
US-09-457-571-12
Query Match 82.1%; Score 23; DB 4; Length 1989;
Best Local Similarity 100.0%; Pred. No. 7,2e-14; Indels 0; Gaps 0;
Matches 23; Conservative 0; Mismatches 0;
CY 6 MMVVGIVLVNLFLLILSSSF 28
DB 949 MMVVGIVLVNLFLLILSSSF 971

RESULT 21
US-10-162-012-24
Sequence 24, Application US/10162012
GENERAL INFORMATION:
APPLICANT: Curtiss, Rony A.J.
APPLICANT: Siles-Santiago, Inmaculada
TITLE OF INVENTION: NOVEL HUMAN ION CHANNEL AND TRANSPORTER FAMILY MEMBERS
FILE REFERENCE: 10448-190001
CURRENT FILING DATE: 2002-06-04
PRIOR APPLICATION NUMBER: US 60/209,845
PRIOR FILING DATE: 2000-06-06-06-09/875,321
PRIOR APPLICATION NUMBER: US 09/875,321
PRIOR FILING DATE: 2001-06-06
PRIOR APPLICATION NUMBER: PCT/US01/18340
PRIOR FILING DATE: 2001-06-06
PRIOR APPLICATION NUMBER: US 60/209,257
PRIOR FILING DATE: 2000-06-05
PRIOR APPLICATION NUMBER: US 09/875,423
PRIOR FILING DATE: 2001-06-05
PRIOR APPLICATION NUMBER: PCT/US01/18398
PRIOR FILING DATE: 2001-06-05
PRIOR APPLICATION NUMBER: US 60/209,238
PRIOR FILING DATE: 2000-06-05
PRIOR APPLICATION NUMBER: US 09/875,363
PRIOR FILING DATE: 2001-06-05
PRIOR APPLICATION NUMBER: PCT/US01/18247
PRIOR FILING DATE: 2001-06-05
PRIOR APPLICATION NUMBER: US 60/227,068
PRIOR FILING DATE: 2000-08-22
PRIOR APPLICATION NUMBER: US 09/928,530
PRIOR FILING DATE: 2001-08-13
PRIOR APPLICATION NUMBER: PCT/US01/25475
PRIOR FILING DATE: 2001-08-15
PRIOR APPLICATION NUMBER: US 60/226,770
PRIOR FILING DATE: 2000-08-21
PRIOR APPLICATION NUMBER: US 09/934,421
PRIOR FILING DATE: 2001-08-21
PRIOR APPLICATION NUMBER: PCT/US01/26096
PRIOR FILING DATE: 2001-08-21
PRIOR APPLICATION NUMBER: US 60/279,281
PRIOR FILING DATE: 2001-03-28
PRIOR APPLICATION NUMBER: US 10/109,029

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PRIOR FILING DATE: 2002-03-28
PRIOR APPLICATION NUMBER: PCT/US02/09728
PRIOR FILING DATE: 2002-03-28
PRIOR APPLICATION NUMBER: US 60/290,288
PRIOR FILING DATE: 2001-05-11
PRIOR APPLICATION NUMBER: US (not assigned)
PRIOR FILING DATE: 2002-05-13
PRIOR APPLICATION NUMBER: 4.0
NUMBER OF SEQ ID NOS: 48
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 24
LENGTH: 1836
TYPE: PRT
ORGANISM: Homo sapiens
US-10-162-012-24
Query Match 78.6%; Score 22; DB 4; Length 1836;
Best Local Similarity 100.0%; Pred. No. 6.1e-13; Indels 0; Gaps 0;
Matches 22; Conservative 0; Mismatches 0;
CY 7 MMVVGIVLVNLFLLILSSSF 28
DB 784 MMVVGIVLVNLFLLILSSSF 805

RESULT 22
US-09-562-737-81
Sequence 81, Application US/09562737
GENERAL INFORMATION:
APPLICANT: Herz, Joachim
APPLICANT: Gotthardt, Michael
TITLE OF INVENTION: LDL Receptor Signaling Pathways
FILE REFERENCE: UTSW0708
CURRENT FILING DATE: US/09/562,737
CURRENT FILING DATE: 2000-05-01
NUMBER OF SEQ ID NOS: 132
SOFTWARE: PatentIn Ver. 2.1
SEQ ID NO 81
LENGTH: 1024
TYPE: PRT
ORGANISM: Artificial Sequence
FEATURE: Description of Artificial Sequence: Synthetic
OTHER INFORMATION: Sequence
US-09-562-737-81
Query Match 75.0%; Score 21; DB 4; Length 1024;
Best Local Similarity 100.0%; Pred. No. 3.2e-12; Indels 0; Gaps 0;
Matches 21; Conservative 0; Mismatches 0;
CY 8 MMVVGIVLVNLFLLILSSSF 28
DB 918 MMVVGIVLVNLFLLILSSSF 938

RESULT 23
US-09-634-920-4
Sequence 4, Application US/09634920
GENERAL INFORMATION:
APPLICANT: Splawski, Igor
APPLICANT: Keating, Mark T.
TITLE OF INVENTION: SCNSA AND METHODS FOR DETECTING
FILE REFERENCE: 2323-155
CURRENT FILING DATE: US/09/634,920
CURRENT FILING DATE: 2000-08-09
PRIOR APPLICATION NUMBER: 60/147,488
PRIOR FILING DATE: 2000-03-17
PRIOR APPLICATION NUMBER: 60/147,488
PRIOR FILING DATE: 1999-08-09
NUMBER OF SEQ ID NOS: 4
SOFTWARE: PatentIn Ver. 2

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SEQ ID NO 4
LENGTH: 2016
TYPE: PRT
ORGANISM: Homo sapiens
US-09-634-920-4

Query Match 75.0%; Score 21; DB 3; Length 2016;
Best Local Similarity 100.0%; Pred. No. 6,1e-12;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 VMVIGNLVNLFLLALLSSF 28
Db 922 VMVIGNLVNLFLLALLSSF 942

RESULT 24
US-09-514-907A-2
Sequence 2, Application US/09514907A
Patent No. 6567705
GENERAL INFORMATION:

APPLICANT: Kenneth B. Stokes
TITLE OF INVENTION: SYSTEMS FOR ENHANCING CARDIAC SIGNAL
SENSING BY CARDIAC PACEMAKERS THROUGH
GENETIC TREATMENT
NUMBER OF SEQUENCES: 12
CORRESPONDENCE ADDRESS:
ADDRESS: Woodcock Washburn Kurtz Mackiewicz & No. 6567705stis LLP
STREET: One Liberty Place - 46th Floor
CITY: Philadelphia
STATE: PA
COUNTRY: U.S.A.
ZIP: 19103

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: WordPerfect 6.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/514,907A
FILING DATE: 08-Feb-2000
CLASSIFICATION: <Unknown>

ATTORNEY/AGENT INFORMATION:
NAME: Paul K. Legaard
REGISTRATION NUMBER: 38,534
REFERENCE/DOCKET NUMBER: MEDT-0029/P-3586
TELEPHONE: (215) 568-3100
TELEFAX: (215) 568-3439
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 2016 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: unknown

US-09-514-907A-2
SEQUENCE DESCRIPTION: SEQ ID NO: 2:

Query Match 75.0%; Score 21; DB 4; Length 2016;
Best Local Similarity 100.0%; Pred. No. 6,1e-12;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 VMVIGNLVNLFLLALLSSF 28
Db 922 VMVIGNLVNLFLLALLSSF 942

RESULT 25
US-09-896-994-2
Sequence 2, Application US/09896994
Patent No. 6665563
GENERAL INFORMATION:
APPLICANT: Ken Stokes

Joe e Morissette
TITLE OF INVENTION: SYSTEMS AND METHODS FOR ENHANCING CARDIAC
SIGNAL SENSING BY CARDIAC PACEMAKERS THROUGH GENETIC TREATM
NUMBER OF SEQUENCES: 12
CORRESPONDENCE ADDRESS:
ADDRESS: Woodcock Washburn Kurtz Mackiewicz and No. 6665563ris
STREET: One Liberty Place - 46th Floor
CITY: Philadelphia
STATE: PA
COUNTRY: U.S.A.
ZIP: 19103

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: WordPerfect 6.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/896,994
FILING DATE: 02-Jul-2001
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 09/514,907
FILING DATE: <Unknown>

ATTORNEY/AGENT INFORMATION:
NAME: Paul K. Legaard
REGISTRATION NUMBER: 38,534
REFERENCE/DOCKET NUMBER: MEDT-0029/P-3586
TELEPHONE: (215) 568-3100
TELEFAX: (215) 568-3439
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 2016 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: unknown

US-09-896-994-2
SEQUENCE DESCRIPTION: SEQ ID NO: 2:

Query Match 75.0%; Score 21; DB 4; Length 2016;
Best Local Similarity 100.0%; Pred. No. 6,1e-12;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Search completed: January 27, 2005, 17:54:18
Job time : 23.5 secs

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OM protein - protein search, using bw model

Run on: January 27, 2005, 17:33:04 ; Search time 86.5 Seconds

(without alignments)
116.120 Million cell updates/sec

Title: US-10-608-584-19

Perfect score: 28
Sequence: 1 MYLYFVIFIFGSEFTLNLFIGVIIDNF 28Scoring table: OLIGO
Gapop 60.0 , Gapext 60.0

Searched: 2002273 seqs, 358729299 residues

Word size : 0
Total number of hits satisfying chosen parameters: 2002273Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Listing first 100 summaries

Database : A Geneseq_23Sep04:*

1: geneseqp1980s:*\n2: geneseqp1990s:*\n3: geneseqp2000s:*\n4: geneseqp2001s:*\n5: geneseqp2002s:*\n6: geneseqp2003as:*\n7: geneseqp2003bs:*\n8: geneseqp2004s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	28	100.0	280	8	ADP45244 Human scd
2	28	100.0	1795	7	ADB78596 Human scd
3	28	100.0	1836	7	ADP57388 Human Pro
4	28	100.0	1836	7	ADP59630 Human Pro
5	28	100.0	1836	7	ADP63029 Human Pro
6	28	100.0	1836	8	ADQ17412 Human scd
7	28	100.0	1855	7	ADB78597 Human scd
8	28	100.0	1950	7	ADB78607 Human scd
9	28	100.0	1951	4	AA899678 Human adu
10	28	100.0	1951	4	AA899679 Human neo
11	28	100.0	1951	7	ADP59628 Rat Prote
12	28	100.0	1951	8	ADL06576 Human tum
13	28	100.0	1962	5	AAE20511 Human ion
14	28	100.0	1973	5	AAE20516 Human ion
15	28	100.0	1981	5	AAE20515 Human SCN
16	28	100.0	1981	5	AAE20516 Human SCN
17	28	100.0	1998	5	AAE20510 Human ion
18	28	100.0	1998	5	AAE20510 Human ion
19	28	100.0	1999	5	AAE20510 Human ion
20	28	100.0	2000	8	AAE20510 Human ion
21	28	100.0	2000	8	AAE20510 Human ion
22	28	100.0	2005	4	AA899676 Human adu
23	28	100.0	2005	4	AA899677 Human neo
24	28	100.0	2005	5	AA899677 Human neo
25	28	100.0	2005	7	ADB78604 Human scd

26	28	100.0	2005	7	ADB78605 Human scd
27	28	100.0	2005	7	ADP46947 Human SCN
28	28	100.0	2009	4	AA899674 Human adu
29	28	100.0	2009	5	AAE20515 Human ion
30	28	100.0	2009	5	AAE20515 Human ion
31	28	100.0	2009	5	AAE20515 Human ion
32	28	100.0	2009	5	AAE20515 Human ion
33	28	100.0	2009	5	AAE20515 Human ion
34	28	100.0	2009	5	AAE20515 Human ion
35	28	100.0	2009	5	AAE20515 Human ion
36	28	100.0	2009	5	AAE20515 Human ion
37	28	100.0	2009	5	AAE20515 Human ion
38	28	100.0	2009	5	AAE20515 Human ion
39	28	100.0	2009	5	AAE20515 Human ion
40	28	100.0	2009	5	AAE20515 Human ion
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42	28	100.0	2009	5	AAE20515 Human ion
43	28	100.0	2009	5	AAE20515 Human ion
44	28	100.0	2009	5	AAE20515 Human ion
45	28	100.0	2009	5	AAE20515 Human ion
46	28	100.0	2009	5	AAE20515 Human ion
47	28	100.0	2009	5	AAE20515 Human ion
48	28	100.0	2009	5	AAE20515 Human ion
49	28	100.0	2009	5	AAE20515 Human ion
50	28	100.0	2009	5	AAE20515 Human ion
51	28	100.0	2009	5	AAE20515 Human ion
52	28	100.0	2009	5	AAE20515 Human ion
53	28	100.0	2009	5	AAE20515 Human ion
54	28	100.0	2009	5	AAE20515 Human ion
55	28	100.0	2009	5	AAE20515 Human ion
56	28	100.0	2009	5	AAE20515 Human ion
57	28	100.0	2009	5	AAE20515 Human ion
58	28	100.0	2009	5	AAE20515 Human ion
59	28	100.0	2009	5	AAE20515 Human ion
60	28	100.0	2009	5	AAE20515 Human ion
61	28	100.0	2009	5	AAE20515 Human ion
62	28	100.0	2009	5	AAE20515 Human ion
63	28	100.0	2009	5	AAE20515 Human ion
64	28	100.0	2009	5	AAE20515 Human ion
65	28	100.0	2009	5	AAE20515 Human ion
66	28	100.0	2009	5	AAE20515 Human ion
67	28	100.0	2009	5	AAE20515 Human ion
68	28	100.0	2009	5	AAE20515 Human ion
69	28	100.0	2009	5	AAE20515 Human ion
70	28	100.0	2009	5	AAE20515 Human ion
71	28	100.0	2009	5	AAE20515 Human ion
72	28	100.0	2009	5	AAE20515 Human ion
73	28	100.0	2009	5	AAE20515 Human ion
74	28	100.0	2009	5	AAE20515 Human ion
75	28	100.0	2009	5	AAE20515 Human ion
76	28	100.0	2009	5	AAE20515 Human ion
77	28	100.0	2009	5	AAE20515 Human ion
78	28	100.0	2009	5	AAE20515 Human ion
79	28	100.0	2009	5	AAE20515 Human ion
80	28	100.0	2009	5	AAE20515 Human ion
81	28	100.0	2009	5	AAE20515 Human ion
82	28	100.0	2009	5	AAE20515 Human ion
83	28	100.0	2009	5	AAE20515 Human ion
84	28	100.0	2009	5	AAE20515 Human ion
85	28	100.0	2009	5	AAE20515 Human ion
86	28	100.0	2009	5	AAE20515 Human ion
87	28	100.0	2009	5	AAE20515 Human ion
88	28	100.0	2009	5	AAE20515 Human ion
89	28	100.0	2009	5	AAE20515 Human ion
90	28	100.0	2009	5	AAE20515 Human ion
91	28	100.0	2009	5	AAE20515 Human ion
92	28	100.0	2009	5	AAE20515 Human ion
93	28	100.0	2009	5	AAE20515 Human ion
94	28	100.0	2009	5	AAE20515 Human ion
95	28	100.0	2009	5	AAE20515 Human ion
96	28	100.0	2009	5	AAE20515 Human ion
97	28	100.0	2009	5	AAE20515 Human ion
98	28	100.0	2009	5	AAE20515 Human ion

99 21 75.0 1978 7 ADE54549
100 21 75.0 1989 2 AAR99640

ADe54549 Human Pro
Aar99640 Periphera

ALIGNMENTS

RESULT 1
ID ADF45244 standard; protein; 280 AA.

XX ADF45244;

XX 26-FEB-2004 (first entry)

DE Human sodium channel alpha subunit amino acid sequence.

XX human; cation channel; INPIONCH05; INPIONCH06; tetrameric cation channel;
XX antiinfectivity; neuroprotective; cardiovascular; immunosuppressive;
XX cerebroprotective; vasotropic; contraceptive; vaccine; infertility;
XX neurological disorder; cardiovascular disorder; autoimmune disease;
XX stroke; stroke-related disorder; pathological condition; calcium channel.

OS Homo sapiens.

XX WO2003099865-A1.

XX 04-DEC-2003.

XX 23-MAY-2003; 2003WO-GB002270.

XX 24-MAY-2002; 2002GB-00012067.

XX (INPH-) INPHARMATICA LTD.

XX Lobley AE, Michalovich D, Allen KE, Reynolds L, Pierron VN;

XX WPI; 2004-053233/05.

XX New INPIONCH05 and INPIONCH06 polypeptides, useful as a contraceptive
XX agent, or for diagnosing and treating a disease or disorder, e.g.
XX infertility, neurological disorder, cardiovascular disorder, autoimmune
XX disease or stroke.

XX Example 1; Fig 2; 96pp; English.

XX The present invention describes cation channel polypeptides (I), termed
XX INPIONCH05 and INPIONCH06. Also described: (1) a purified nucleic acid
XX molecule (II), which: (a) encodes (I); or (b) hybridizes under high
XX stringency conditions with the nucleic acid molecule of (a); (2) a vector
XX comprising (II); (3) a host cell transformed with the vector; (4) a
XX tetrameric cation channel comprising (I); (5) a ligand which binds
XX specifically to, and which preferably inhibits the activity of (I) or the
XX tetrameric cation channel; (6) a compound that either increases or
XX decreases the level of expression or activity of (I) or the tetrameric
XX cation channel; (7) a method of diagnosing a disease in a patient; (8) a
XX pharmaceutical composition comprising (I), (II), vector, host cell,
XX comprising (I) or (II); (10) a method of treating a disease in a patient;
XX (11) a method of monitoring the therapeutic treatment of disease in a
XX patient; (12) a method for the identification of a compound that is
XX an effective in the treatment and/or diagnosis of a disease/disorder; (13)
XX an array of nucleic acid molecules, at least one of which is (II), or one
XX or more antibodies that bind to (I) or to the tetrameric cation channel,
XX and a reagent useful for the detection of a binding reaction between the
XX antibody and (I) or tetrameric cation channel; (14) a transgenic or
XX knockout non-human animal that has been transformed to express higher,
XX lower or absent levels of (I); and (15) a method for screening for a
XX compound effective to treat a disease or disorder. (I) has
XX antiinfectivity, neuroprotective, cardiovascular, immunosuppressive,
XX cerebroprotective, vasotropic and contraceptive activities, and can be
XX used in vaccines, (I), (II), vectors, tetrameric cation channels,
XX ligands, compounds, or compositions of the present invention may be used

CC as contraceptive agents, preferably a non-hormonal contraceptive agent,
CC in therapy or diagnosis of a disease or disorder, or in manufacturing a
CC medicament for the treatment of a disease or disorder, e.g. infertility,
CC neurological disorder, cardiovascular disorder, autoimmune disease,
CC stroke, stroke-related disorders or other pathological condition. (I) is
CC useful as a cation channel or a calcium channel. (I) or the tetrameric
CC cation channel is useful as a sperm-specific CatSper channel. The present
CC sequence is used in the exemplification of the present invention.

XX SQ Sequence 280 AA;

Query Match 100.0%; Score 28; DB 8; Length 280;
Best Local Similarity 100.0%; Pred. No. 2,4e-21;

Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MLYFVILIFGSPFTNLFGVITIDNF 28
DB 246 MLYFVILIFGSPFTNLFGVITIDNF 273

RESULT 2
ID ADB78596 standard; protein; 1795 AA.

XX ADB78596;

XX 04-DEC-2003 (first entry)

XX Human sodium channel subunit mutant SEQ ID NO:140.

XX mutein; mutant; ion channel; ion channel subunit; ICS; nootropic;
XX neuroprotective; inotropic; antipyretic; antiarrhythmic; antimigraine;
XX antidepressant; antiparkinsonian; neuroleptic; tranquilizer; analgesic;
XX nephrotropic; antidiabetic; ophthalmological; epilepsy;
XX ion channel dysfunction; human.

XX Synthetic.

XX OS Homo sapiens.

XX WO2003008574-A1.

XX 30-JAN-2003.

XX 08-JUL-2002; 2002WO-AU000910.

XX 18-JUL-2001; 2001AU-00006452.

XX 05-MAR-2002; 2002AU-00000910.

XX 13-MAY-2002; 2002AU-00002292.

XX (BION-) BIONOMICS LTD.

XX (WALL-) WALLACE R W.

XX Muller JC, Harkin LA, Dibbens LM, Phillips HA, Heron SE;

XX Berkovic SF, Scheffer IE;

XX WPI; 2003-239332/23.

XX N-PSDB; ADB78635.

XX Identifying predisposition to an ion channel dysfunction, such as

XX periodic paralysis, cardiac arrhythmias, migraine, Alzheimer's disease,

XX schizophrenia, anxiety and depression, by detecting encoding-gene

XX mutation events.

XX Claim 13; SEQ ID NO 140; 106pp; English.

XX The invention relates to a novel method for identifying a subject
XX predisposed to a disorder associated with ion channel dysfunction. The
XX method comprises ascertaining if at least one of the genes encoding ion
XX channel subunits (ICS) has undergone a mutation event so that a cDNA
XX derived from the subject has any of 134 nucleotide sequences. The method
XX of the invention has nootropic, neuroprotective, inotropic, antipyretic,
XX antiarrhythmic, antimigraine, antidepressant, antiparkinsonian,
XX neuroleptic, tranquilizer, analgesic, nephrotropic, antidiabetic, and

ophthalmological activity. A polynucleotide of the invention acts as an ion channel agonist, or ion channel antagonist. The methods, isolated nucleic acids, polypeptides, antibody, selective agonist, antagonist or modulator of an ion channel, cells and genetically modified non-human animal, are useful for the diagnosis and treatment of epilepsy and/or a disorder associated with ion channel dysfunction, such as hyper- or hypokalemic periodic paralysis, myotonia, malignant hyperthermia, myasthenia, cardiac arrhythmias, episodic ataxia, migraine, Alzheimer's disease, Parkinson's disease, schizophrenia, hyperkplexia, anxiety, depression, phobic obsessive symptoms, neuropathic pain, inflammatory pain, chronic/acute pain, Bartter's syndrome, polycystic kidney disease, Dent's disease, hyperinsulinaemic hypoglycaemia of infancy, cystic fibrosis, congenital stationary night blindness and total colour blindness. The present sequence represents a mutant protein of the invention. The sequence data for this patent is not represented in the printed specification, but was obtained in electronic format directly from WIPO at ftp.wipo.int/pub/published_pat_sequences.

Sequence 1795 AA;

Query Match 100.0%; Score 28; DB 7; Length 1795;
Best Local Similarity 100.0%; Pred. No. 1.3e-20;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 MYLVFVFIIFGSPFTLNLFIGVIIDNF 28
1459 MYLVFVFIIFGSPFTLNLFIGVIIDNF 1486

RESULT 3
ADE57388 ID ADE57388 standard; protein; 1836 AA.

AC ADE57388;

DT 29-JAN-2004 (first entry)

DE Human Protein XP_008249, SEQ ID NO 3249.

Human; pain; neuronal tissue; gene therapy;
spinal segmental nerve injury; chronic constriction injury; CCI;
spared nerve injury; SN1; Chung.

OS Homo sapiens.

PN WO2003016475-A2.

PD 27-FEB-2003.

PF 14-AUG-2002; 2002MO-US025765.

PR 14-AUG-2001; 2001US-0312147P.

PR 01-NOV-2001; 2001US-0346382P.

PR 26-NOV-2001; 2001US-0333347P.

PA (GENO) GEN HOSPITAL CORP.

PA (FARB) BAYER AG.

PI Woolf C, D'urso D, Befort K, Costigan M;

DR WPI; 2003-268312/26.

DR GENBANK; XP_008249.

PT New composition comprising two or more isolated polypeptides, useful for

PT preparing a medicament for treating pain in an animal.

XX Claim 1; Page; 1017pp; English.

CC The invention discloses a composition comprising two or more isolated rat

CC or human polynucleotides or a polynucleotide which represents a fragment,

CC derivative or allelic variation of the nucleic acid sequence. Also

CC claimed are a vector comprising the novel polynucleotide, a host cell

CC comprising the vector, a method for identifying a nucleotide sequence

CC which is differentially regulated in an animal subjected to pain and a
CC kit to perform the method, an array, a method for identifying an agent
CC that increases or decreases the expression of the polynucleotide sequence
CC that is differentially expressed in neuronal tissue of a first animal
CC subjected to pain, a method for identifying a compound which regulates
CC the expression of a polynucleotide sequence which is differentially
CC expressed in an animal subjected to pain, a method for identifying a
CC compound that regulates the activity of one or more of the
CC polynucleotides, a method for producing a pharmaceutical composition, a
CC method for identifying a compound or small molecule that regulates the
CC activity in an animal of one or more of the polypeptides given in the
CC specification, a method for identifying a compound useful in treating
CC pain and a pharmaceutical composition comprising the one or more
CC polypeptides or their antibodies. The polynucleotide or the compound that
CC modulates its activity is useful for preparing a medicament for treating
CC pain (e.g. spinal segmental nerve injury (Chung), chronic constriction
CC injury (CCI) and spared nerve injury (SN1)) in an animal (e.g. gene
CC therapy). The sequence presented is a human protein (shown in Table 2 of
CC the specification) which is differentially expressed during pain. Note:
CC The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic form directly from WIPO at
CC ftp.wipo.int/pub/published_pat_sequences.

Sequence 1836 AA;

Query Match 100.0%; Score 28; DB 7; Length 1836;
Best Local Similarity 100.0%; Pred. No. 1.3e-20;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 MYLVFVFIIFGSPFTLNLFIGVIIDNF 28
1271 MYLVFVFIIFGSPFTLNLFIGVIIDNF 1298

RESULT 4
ADE59630 ID ADE59630 standard; protein; 1836 AA.

AC ADE59630;

DT 29-JAN-2004 (first entry)

DE Human Protein XP_008249, SEQ ID NO 5526.

Human; pain; neuronal tissue; gene therapy;
spinal segmental nerve injury; chronic constriction injury; CCI;
spared nerve injury; SN1; Chung.

OS Homo sapiens.

PN WO2003016475-A2.

PD 27-FEB-2003.

PF 14-AUG-2002; 2002MO-US025765.

PR 14-AUG-2001; 2001US-0312147P.

PR 01-NOV-2001; 2001US-0346382P.

PR 26-NOV-2001; 2001US-0333347P.

PA (GENO) GEN HOSPITAL CORP.

PA (FARB) BAYER AG.

PI Woolf C, D'urso D, Befort K, Costigan M;

DR WPI; 2003-268312/26.

DR GENBANK; XP_008249.

PT New composition comprising two or more isolated polypeptides, useful for

PT preparing a medicament for treating pain in an animal.

XX Claim 1; Page; 1017pp; English.

CC The invention discloses a composition comprising two or more isolated rat

CC or human polynucleotides or a polynucleotide which represents a fragment,

CC derivative or allelic variation of the nucleic acid sequence. Also

CC The invention discloses a composition comprising two or more isolated rat
 CC or human polynucleotides or a polynucleotide which represents a fragment,
 CC derivative or allelic variation of the nucleic acid sequence. Also
 CC claimed are a vector comprising the novel polynucleotide, a host cell
 CC comprising the vector, a method for identifying a nucleotide sequence
 CC which is differentially regulated in an animal subjected to pain and a
 CC kit to perform the method, an array, a method for identifying an agent
 CC that increases or decreases the expression of the polynucleotide sequence
 CC that is differentially expressed in neuronal tissue of a first animal
 CC subjected to pain, a method for identifying a compound which regulates
 CC the expression of a polynucleotide sequence which is differentially
 CC expressed in an animal subjected to pain, a method for identifying a
 CC compound that regulates the activity of one or more of the
 CC polynucleotides, a method for producing a pharmaceutical composition, a
 CC method for identifying a compound or small molecule that regulates the
 CC activity in an animal of one or more of the polypeptides given in the
 CC pain and a pharmaceutical composition comprising the one or more
 CC polypeptides or their antibodies. The polynucleotide or the compound that
 CC pain (e.g. spinal segmental nerve injury (Chung), chronic constriction
 CC injury (CCI) and spared nerve injury (SNI)) in an animal (e.g. gene
 CC therapy). The sequence presented is a human protein (shown in Table 2 of
 CC the specification) which is differentially expressed during pain. Note:
 CC The sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic form directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences.

SO Sequence 1836 AA;

Query Match 100.0%; Score 28; DB 7; Length 1836;
 Best Local Similarity 100.0%; Pred. No. 1.3e-20;

Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 MYLFFVFIIFGSEFTNLFIVGIIDNF 28
 |||||
 DB 1271 MYLFFVFIIFGSEFTNLFIVGIIDNF 1298

RESULT 5
 ADE63029
 ID ADE63029 standard; protein; 1836 AA.
 XX
 AC ADE63029;

DT 29-JAN-2004 (first entry)
 XX
 DE Human Protein XP_008249, SEQ ID NO 8963.

XX Human; pain; neuronal tissue; gene therapy;
 KM spinal segmental nerve injury; chronic constriction injury; CCI,
 XX spared nerve injury; SNI; Chung.

OS Homo sapiens.

XX WO2003016475-A2.

PD 27-FEB-2003.

PF 14-AUG-2002; 2002WO-US025765.

PR 14-AUG-2001; 2001US-0312147P.

PR 01-NOV-2001; 2001US-0346383P.

PR 26-NOV-2001; 2001US-0333347P.

PA (GENO) GEN HOSPITAL CORP.

PA (FARB) BAYER AG.

PI Woolf C, D'urso D, Befort K, Costigan M;
 DR WPI; 2003-268312/26.
 DR GENBANK; XP_008249.
 XX

PT New composition comprising two or more isolated polypeptides, useful for
 PT preparing a medicament for treating pain in an animal.

PS Claim 1; Page; 1017pp; English.

CC The invention discloses a composition comprising two or more isolated rat
 CC or human polynucleotides or a polynucleotide which represents a fragment,
 CC derivative or allelic variation of the nucleic acid sequence. Also
 CC claimed are a vector comprising the novel polynucleotide, a host cell
 CC comprising the vector, a method for identifying a nucleotide sequence
 CC which is differentially regulated in an animal subjected to pain and a
 CC kit to perform the method, an array, a method for identifying an agent
 CC that increases or decreases the expression of the polynucleotide sequence
 CC that is differentially expressed in neuronal tissue of a first animal
 CC subjected to pain, a method for identifying a compound which regulates
 CC the expression of a polynucleotide sequence which is differentially
 CC expressed in an animal subjected to pain, a method for identifying a
 CC compound that regulates the activity of one or more of the
 CC polynucleotides, a method for producing a pharmaceutical composition, a
 CC method for identifying a compound or small molecule that regulates the
 CC activity in an animal of one or more of the polypeptides given in the
 CC pain and a pharmaceutical composition comprising the one or more
 CC polypeptides or their antibodies. The polynucleotide or the compound that
 CC pain (e.g. spinal segmental nerve injury (Chung), chronic constriction
 CC injury (CCI) and spared nerve injury (SNI)) in an animal (e.g. gene
 CC therapy). The sequence presented is a human protein (shown in Table 2 of
 CC the specification) which is differentially expressed during pain. Note:
 CC The sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic form directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences.

SO Sequence 1836 AA;

Query Match 100.0%; Score 28; DB 7; Length 1836;
 Best Local Similarity 100.0%; Pred. No. 1.3e-20;

Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 MYLFFVFIIFGSEFTNLFIVGIIDNF 28
 |||||
 DB 1271 MYLFFVFIIFGSEFTNLFIVGIIDNF 1298

RESULT 6
 ADQ17412
 ID ADQ17412 standard; protein; 1836 AA.
 XX
 AC ADQ17412;

DT 26-AUG-2004 (first entry)

XX Human soft tissue sarcoma-upregulated protein - SEQ ID 229.

XX soft tissue sarcoma; cytostatic; gene therapy; vaccine; screening; human.

OS Homo sapiens.

XX WO2004048938-A2.

PD 10-JUN-2004.

PF 26-NOV-2003; 2003WO-US038193.

PR 26-NOV-2002; 2002US-0429739P.

PA (PROT-) PROTEIN DESIGN LABS INC.

PA Aziz N, Ginsburg WM, Zlotnick A;

DR WPI; 2004-441208/41.

PT Early detection of soft tissue sarcoma comprises determining expression

PI Berkovic SF, Scheffer IE;
 XX WPI; 2003-239332/23.
 DR N-PSDB; ADB78646.
 XX
 PT Identifying predisposition to an ion channel dysfunction, such as
 PT periodic paralysis, cardiac arrhythmias, migraine, Alzheimer's disease,
 PT schizophrenia, anxiety and depression, by detecting encoding-gene
 PT mutation events.
 XX
 PS Claim 13; SEQ ID NO 151; 106pp; English.
 CC
 CC The invention relates to a novel method for identifying a subject
 CC predisposed to a disorder associated with ion channel dysfunction. The
 CC method comprises ascertaining if at least one of the genes encoding ion
 CC channel subunits (ICs) has undergone a mutation event so that a cDNA
 CC derived from the subject has any of 134 nucleotide sequences. The method
 CC of the invention has nucleotide, neuroprotective, isotropic, antipyretic,
 CC antihypertensive, antimigraine, antidepressant, antiparkinsonian,
 CC neuroleptic, tranquiliser, analgesic, nephrotoxic, antidiabetic, and
 CC ion channel agonist, or ion channel antagonist. The methods, isolated
 CC modulator of an ion channel, cells and genetically modified non-human
 CC animal, are useful for the diagnosis and treatment of epilepsy and/or a
 CC disorder associated with ion channel dysfunction, such as hyper- or hypo-
 CC kalemia, cardiac arrhythmias, myotonias, malignant hyperthermia,
 CC depression, Parkinson's disease, schizophrenia, hyperkplexia, anxiety,
 CC pain, chronic/acute pain, Bartter's syndrome, polycystic kidney disease,
 CC Dent's disease, hyperinsulinaemic hypoglycaemia of infancy, cystic
 CC fibrosis, congenital stationary night blindness and total colour
 CC blindness. The present sequence data for this patent is not represented in the
 CC printed specification, but was obtained in electronic format directly
 CC from WIPO at ftp.wipo.int/pub/published_pat_sequences.
 XX
 SQ Sequence 1950 AA;
 Query Match 100.0%; Score 28; DB 7; Length 1950;
 Best Local Similarity 100.0%; Pred. No. 1.4e-20;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 MYLFFVFIIFGSGFTNLFGVITDNF 28
 DB 1394 MYLFFVFIIFGSGFTNLFGVITDNF 1421
 RESULT 9
 AAB99678
 ID AAB99678 standard; protein; 1951 AA.
 AC AAB99678;
 XX
 DT 04-SEP-2001 (first entry)
 XX
 DE Human adult form of SCN3A protein sequence SEQ ID NO:67.
 XX
 KW Human; epilepsy; chromosome 2; SCN1A; SCN2A; SCN3A; identification;
 KW diagnosis; mutation; chromosome 2q23-q31; neurological disorder;
 XX anticonvulsant; neuroprotective.
 OS Homo sapiens.
 XX
 PN WO200138564-A2.
 PD 31-MAY-2001.
 XX
 PF 24-NOV-2000; 2000WO-CA001404.
 XX
 PR 26-NOV-1999; 99US-0167623P.
 XX

PA (UTMC-) UNIV MCGILL.
 XX
 PI Rouleau GA, Lafreniere RG, Rochefort D, Cossette P, Ragsdale D;
 XX WPI; 2001-355945/37.
 DR N-PSDB; AAH55823.
 XX
 PT Determining a predisposition to epilepsy and/or development of epilepsy
 PT comprises determining the genotype of SCN1A, SCN2A and/or SCN3A, or a DNA
 PT variant, equivalent, or mutation which shows a linkage disequilibrium.
 XX
 PS Disclosure; Page 157-165; 268pp; English.
 CC
 CC The present invention describes a method (M1) of determining an
 CC individual's predisposition to epilepsy and/or development of epilepsy,
 CC as well as predicting the individual's response to medication. The method
 CC comprises determining the genotype of at least one gene selected from
 CC SCN1A, SCN2A or SCN3A, or a DNA variant, equivalent or mutation which
 CC shows a linkage disequilibrium. SCN1A, SCN2A and SCN3A are all sodium
 CC channel genes located on chromosome 2. The idiopathic generalised
 CC epilepsy (IGE) gene is more specifically localised on chromosome 2q23-
 CC q31. Compounds identified as modulators of the biological activity of
 CC SCN1A, SCN2A or SCN3A proteins or genes, are useful for treating epilepsy
 CC or other neurological disorders. They have anticonvulsant and
 CC neuroprotective activities. AAH55763 to AAH56164 and AAB99674 to AAB99679
 CC represent SCN1A, SCN2A, and SCN3A cDNAs, gene fragments, PCR primers,
 CC oligonucleotides and proteins given in the exemplification of the present
 CC invention
 XX
 SQ Sequence 1951 AA;
 Query Match 100.0%; Score 28; DB 4; Length 1951;
 Best Local Similarity 100.0%; Pred. No. 1.4e-20;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 MYLFFVFIIFGSGFTNLFGVITDNF 28
 DB 1395 MYLFFVFIIFGSGFTNLFGVITDNF 1422
 RESULT 10
 AAB99679
 ID AAB99679 standard; protein; 1951 AA.
 AC AAB99679;
 XX
 DT 04-SEP-2001 (first entry)
 XX
 DE Human neonatal form of SCN3A protein sequence SEQ ID NO:68.
 XX
 KW Human; epilepsy; chromosome 2; SCN1A; SCN2A; SCN3A; identification;
 KW diagnosis; mutation; chromosome 2q23-q31; neurological disorder;
 XX anticonvulsant; neuroprotective.
 OS Homo sapiens.
 XX
 PN WO200138564-A2.
 PD 31-MAY-2001.
 XX
 PF 24-NOV-2000; 2000WO-CA001404.
 XX
 PR 26-NOV-1999; 99US-0167623P.
 XX
 PA (UTMC-) UNIV MCGILL.
 XX
 PI Rouleau GA, Lafreniere RG, Rochefort D, Cossette P, Ragsdale D;
 XX WPI; 2001-355945/37.
 DR N-PSDB; AAH55824.
 XX
 PT Determining a predisposition to epilepsy and/or development of epilepsy
 PT comprises determining the genotype of SCN1A, SCN2A and/or SCN3A, or a DNA

PT variant, equivalent, or mutation which shows a linkage disequilibrium.
XX
XX Disclosure; Page 165-172; 268pp; English.
XX
XX The present invention describes a method (M1) of determining an
CC individual's predisposition to epilepsy and/or development of epilepsy,
CC as well as predicting the individual's response to medication. The method
CC comprises determining the genotype of at least one gene selected from
CC SCNA1, SCNA2 or SCNA3, or a DNA variant, equivalent, or mutation which
CC shows a linkage disequilibrium. SCNA1, SCNA2 and SCNA3 are all sodium
CC channel genes located on chromosome 2. The idiopathic generalised
CC epilepsy (IGE) gene is more specifically localised on chromosome 2q23-
CC q31. Compounds identified as modulators of the biological activity of
CC SCNA1, SCNA2 or SCNA3 proteins or genes, are useful for treating epilepsy
CC or other neurological disorders. They have anticonvulsant and
CC neuroprotective activities. AAH5763 to AAH56164 and AAB99674 to AAB99679
CC represent SCNA1, SCNA2, and SCNA3 cDNAs, gene fragments, PCR primers,
CC oligonucleotides and proteins given in the exemplification of the present
CC invention
XX
SQ Sequence 1951 AA;

Query Match 100.0%; Score 28; DB 4; Length 1951;
Best Local Similarity 100.0%; Pred. No. 1.4e-20;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MYLYFVIFPIFGSFFTLNFIGVIIDNF 28
Db 1395 MYLYFVIFPIFGSFFTLNFIGVIIDNF 1422
|||||

RESULT 11
ADES9628
ID ADES9628 standard; protein; 1951 AA.
XX
XX ADES9628;
AC
XX
DT 29-JAN-2004 (first entry)
XX
XX Rat Protein NP_037251, SEQ ID NO 5524.
DE
XX
XX Rat; pain; neuronal tissue; gene therapy; spinal segmental nerve injury;
KW chronic constriction injury; CCI; spared nerve injury; SN1; Chung.
XX
XX Rattus norvegicus.
OS
XX
XX WO2003016475-A2.
PN
XX
XX 27-FEB-2003.
PD
XX
XX 14-AUG-2002; 2002WO-US025765.
PF
XX
XX 14-AUG-2001; 2001US-0312147P.
PR 01-NOV-2001; 2001US-0346382P.
PR 26-NOV-2001; 2001US-0333347P.
XX
XX (GHEO) GEN HOSPITAL CORP.
PA (FARB) BAYER AG.
XX
XX Woolf C, D'urso D, Befort K, Costigan M;
PI
XX WPI, 2003-268312/26.
DR
XX GENBANK; NF_037251.
DR
XX
XX New composition comprising two or more isolated polypeptides, useful for
PT preparing a medicament for treating pain in an animal.
PT
XX
XX Claim 1; Page; 1017pp; English.
PS
XX
XX The invention discloses a composition comprising two or more isolated rat
CC or human polynucleotides or a polynucleotide which represents a fragment,
CC derivative or allelic variation of the nucleic acid sequence. Also
CC claimed are a vector comprising the novel polynucleotide, a host cell

CC comprising the vector, a method for identifying a nucleotide sequence
CC which is differentially regulated in an animal subjected to pain and a
CC kit to perform the method, an array, a method for identifying an agent
CC that increases or decreases the expression of the polynucleotide sequence
CC that is differentially expressed in neuronal tissue of a first animal
CC subjected to pain, a method for identifying a compound which regulates
CC the expression of a polynucleotide sequence which is differentially
CC expressed in an animal subjected to pain, a method for identifying a
CC compound that regulates the activity of one or more of the
CC polynucleotides, a method for producing a pharmaceutical composition, a
CC method for identifying a compound or small molecule that regulates the
CC activity in an animal of one or more of the polypeptides given in the
CC specification, a method for identifying a compound useful in treating
CC pain and a pharmaceutical composition comprising the one or more
CC polypeptides or their antibodies. The polynucleotide or the compound that
CC modulates its activity is useful for preparing a medicament for treating
CC pain (e.g. spinal segmental nerve injury (Chung), chronic constriction
CC injury (CCI) and spared nerve injury (SN1)) in an animal (e.g. gene
CC therapy). The sequence presented is a rat protein (shown in Table 2 of
CC the specification) which is differentially expressed during pain. Note:
CC The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic form directly from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences.
XX
XX
SQ Sequence 1951 AA;

Query Match 100.0%; Score 28; DB 7; Length 1951;
Best Local Similarity 100.0%; Pred. No. 1.4e-20;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MYLYFVIFPIFGSFFTLNFIGVIIDNF 28
Db 1395 MYLYFVIFPIFGSFFTLNFIGVIIDNF 1422
|||||

RESULT 12
ADL06576
ID ADL06576 standard; protein; 1951 AA.
XX
XX ADL06576;
AC
XX
XX 20-MAY-2004 (first entry)
DT
XX
XX Human tumour-associated antigenic target (TAT) polypeptide #75.
DE
XX
XX Human; tumour-associated antigenic target; TAT; cell death; tumour;
KW cancer; cytostatic.
XX
XX Homo sapiens.
OS
XX
XX WO2004016225-A2.
PN
XX
XX 26-FEB-2004.
PD
XX
XX 19-AUG-2003; 2003WO-US025892.
PF
XX
XX 19-AUG-2002; 2002US-0404809P.
PR 21-AUG-2002; 2002US-0405645P.
PR 23-SEP-2002; 2002US-0413192P.
PR 15-OCT-2002; 2002US-0419008P.
PR 15-NOV-2002; 2002US-0426847P.
PR 02-JUL-2003; 2003US-0484959P.
XX
XX (GETH) GENENTECH INC.
PA
XX
XX Desauvage FU, Frantz G, Hillan KJ, Polakis P, Polson A, Smith V,
PI Spencer SD, Wu TD, Zhang Z;
XX
XX WPI, 2004-257144/24.
DR
XX N-PSDB; ADL06499.
DR
XX
XX New antibody that binds to a tumor-associated antigenic target (TAT)
PT polypeptide, useful for preparing a composition for diagnosing or

PT treating cancer.
XX
PS Claim 2; SEQ ID NO 156; 319pp; English.
XX
CC The present invention relates to the isolation of human tumour-associated
CC antigenic target (TAT) polynucleotide and polypeptide sequences. Also
CC disclosed is an antibody that binds to a TAT polypeptide. The antibody is
CC a monoclonal antibody, an antibody fragment, a chimeric antibody or a
CC humanised antibody. It is conjugated to a growth inhibitory agent. It is
CC produced in bacteria or in CHO cells and induces death of a cell to which
CC it binds. The antibody is useful for preparing a composition for
CC diagnosing or treating tumours and cancer. The present sequence
CC represents a human TAT polypeptide of the invention.
SQ
XX
XX Sequence 1951 AA;
SQ
Query Match 100.0%; Score 28; DB 8; Length 1951;
Best Local Similarity 100.0%; Pred. No. 1.4e-20;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 MYLYPVIFIFGSEFTLNLFIGVIIDNF 28
DB 1395 MYLYPVIFIFGSEFTLNLFIGVIIDNF 1422
RESULT 13
AAE20511
ID AAE20511 standard; protein; 1962 AA.
XX
AC AAE20511;
XX
DT 01-JUL-2002 (first entry)
XX
DE Human ion channel protein #2.
XX
DE Human; novel human protein; NHP; voltage-gated sodium channel;
KM gene therapy; bioreactor; mental disorder; biological disorder;
KM medical disorder.
XX
XX Homo sapiens.
OS
FH Key Location/Qualifiers
FT Misc-difference 981 /note= "Encoded by MTG"
FT Misc-difference 1056 /note= "Encoded by RCA"
XX
PN WO200214498-A2.
XX
PD 21-FEB-2002.
XX
PF 15-AUG-2001; 2001WO-US025650.
XX
PR 16-AUG-2000; 2000US-0225989P.
XX
PA (LEXI-) LEXICON GENETICS INC.
XX
PI Turner CA, Mathur B, Mathur D;
XX
DR WPI: 2002-280757/32.
DR N-PSDB; AAD32840.
XX
PT Novel polynucleotides encoding human sodium channel proteins,
PT particularly voltage-gated sodium channel proteins useful for drug
PT screening, diagnosis and in gene therapy of biological disorders.
XX
PS Claim 1; Page 37-41; 83pp; English.
XX
CC The present sequence is novel human protein (NHP), ion channel protein.
CC NHP share structural similarity with mammalian sodium channel proteins
CC particularly voltage-gated sodium channel proteins. NHP oligonucleotides
CC are useful as hybridisation probes for screening libraries and assessing the
CC gene expression patterns. Sequences derived from regions adjacent to the

CC intron/exon boundaries of NHP gene can be used to design primers for use
CC in amplification assays to detect mutations within the exons, splice
CC sites, introns that can be used in diagnostics and pharmacogenomics. NHP
CC nucleotide sequences are useful for drug screening effective in the
CC treatment of symptomatic or phenotypic manifestations of perturbing the
CC normal function of NHP in the body, and nucleotide constructs encoding
CC NHP products are useful to genetically engineer host cells to express NHP
CC products in vivo. These genetically engineered cells function as
CC bioreactors in the body delivering a continuous supply of a NHP, a NHP
CC peptide, or a NHP fusion protein to the body. Nucleotide construct
CC encoding NHP products are also useful in gene therapy for modulating NHP
CC expression and to produce genetically engineered host cells to express
CC NHP products in vivo. NHP nucleotide sequences may also be used as part
CC of ribozyme and/or triplex helix sequences that are useful for NHP gene
CC regulation. The NHP polypeptides are useful for generating antibodies, as
CC reagents in diagnostic assays, for identifying other cellular gene
CC products related to NHP and as reagents in assays for screening for
CC compounds that are useful in the treatment of mental, biological or
CC medical disorders and diseases
SQ
XX
XX Sequence 1962 AA;
SQ
Query Match 100.0%; Score 28; DB 5; Length 1962;
Best Local Similarity 100.0%; Pred. No. 1.4e-20;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 MYLYPVIFIFGSEFTLNLFIGVIIDNF 28
DB 1448 MYLYPVIFIFGSEFTLNLFIGVIIDNF 1475
RESULT 14
AAE20516
ID AAE20516 standard; protein; 1973 AA.
XX
AC AAE20516;
XX
DT 01-JUL-2002 (first entry)
XX
DE Human ion channel protein #7.
XX
DE Human; novel human protein; NHP; voltage-gated sodium channel;
KM gene therapy; bioreactor; mental disorder; biological disorder;
KM medical disorder.
XX
XX Homo sapiens.
OS
FH Key Location/Qualifiers
FT Misc-difference 992 /note= "Encoded by MTG"
FT Misc-difference 1067 /note= "Encoded by RCA"
XX
PN WO200214498-A2.
XX
PD 21-FEB-2002.
XX
PF 15-AUG-2001; 2001WO-US025650.
XX
PR 16-AUG-2000; 2000US-0225989P.
XX
PA (LEXI-) LEXICON GENETICS INC.
XX
PI Turner CA, Mathur B, Mathur D;
XX
DR WPI: 2002-280757/32.
DR N-PSDB; AAD32845.
XX
PT Novel polynucleotides encoding human sodium channel proteins,
PT particularly voltage-gated sodium channel proteins useful for drug
PT screening, diagnosis and in gene therapy of biological disorders.
XX
PS Claim 1; Page 64-68; 83pp; English.

XX The present sequence is novel human protein (NHP), ion channel protein.
CC NHP share structural similarity with mammalian sodium channel proteins.
CC particularly voltage-gated sodium channel proteins. NHP oligonucleotides
CC are useful as hybridisation probes for screening libraries and assessing
CC gene expression patterns. Sequences derived from regions adjacent to the
CC intron/exon boundaries of NHP gene can be used to design primers for use
CC in amplification assays to detect mutations within the exons, splice
CC sites, introns that can be used in diagnostics and pharmacogenomics. NHP
CC nucleotide sequences are useful for drug screening effective in the
CC treatment of symptomatic or phenotypic manifestations of perturbing the
CC normal function of NHP in the body, and nucleotide constructs encoding
CC NHP products are useful to genetically engineer host cells to express NHP
CC products in vivo. These genetically engineered cells function as
CC bioreactors in the body delivering a continuous supply of a NHP, a NHP
CC peptide, or a NHP fusion protein to the body. Nucleotide construct
CC encoding NHP products are also useful in gene therapy for modulating NHP
CC expression and to produce genetically engineered host cells to express
CC NHP products in vivo. NHP nucleotide sequences may also be used as part
CC of ribozyme and/or triple helix sequences that are useful for NHP gene
CC regulation. The NHP polypeptides are useful for generating antibodies, as
CC reagents in diagnostic assays, for identifying other cellular gene
CC products related to NHP and as reagents in assays for screening for
CC compounds that are useful in the treatment of mental, biological or
CC medical disorders and diseases

SQ Sequence 1973 AA:

Query Match 100.0%; Score 28; DB 5; Length 1973;
Best Local Similarity 100.0%; Pred. No. 1.4e-20;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MYLFFVFIIFGSPFTLNLFTGVIIIDNF 28
Db 1459 MYLFFVFIIFGSPFTLNLFTGVIIIDNF 1486

RESULT 15

ID ABR83185 standard; protein; 1981 AA.

XX ABR83185;

DT 15-JAN-2004 (first entry)

DE Human SCN1A splice variant -84P:SCN1ADLP654-681.

XX SCN1A; sodium channel type 1 alpha-subunit; anticonvulsant; analgesic;
KW neuroprotective; anesthetic; cytostatic; cerebroprotective; cardiant;
KW hypotensive; gene therapy; human; splice variant.

XX Homo sapiens.

PN WO2003072751-A2.

PD 04-SEP-2003.

PF 25-FEB-2003; 2003WO-US006010.

PR 25-FEB-2002; 2002US-0359382P.

PA (UVA-) UNIV VANDERBILT.

PI George AL, Loselin C;

XX WPI; 2003-712725/67.

DR N-PSDB; ACF57880.

PT Recombinantly expressed sodium channel type 1 alpha subunit, useful in
screening for modulators, for treating e.g. epilepsy.
PS Disclosure; Page 162-169; 176pp; English.

CC The invention relates to a recombinantly expressed and isolated human
CC SCN1A (sodium channel type 1 alpha-subunit) (I), (II), optionally
CC incorporated into a cell, is used to screen for specific modulators,
CC potentially useful as anticonvulsant, antiepileptic, neuroprotective,
CC analgesic and/or anesthetic agents, e.g. for treating severe myoclonic
CC epilepsy of infancy, stroke, cardiac arrest, hyperkalemic paralysis,
CC motor endplate diseases, hypertension, congestive heart failure and
CC muscular dystrophy also to treat cancer (SCN1A is expressed in prostatic
CC and metastatic cancer cell lines). These activities can also be provided
CC by gene therapy vectors that express (I) or the modulators. The
CC modulators, also antibodies directed against (I), are used to detect
CC sodium channel polypeptides. The present sequence represents a human
CC SCN1A splice variant 84P:SCN1ADLP654-681

SQ Sequence 1981 AA:

Query Match 100.0%; Score 28; DB 7; Length 1981;
Best Local Similarity 100.0%; Pred. No. 1.4e-20;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MYLFFVFIIFGSPFTLNLFTGVIIIDNF 28
Db 1431 MYLFFVFIIFGSPFTLNLFTGVIIIDNF 1458

RESULT 16

ID AAE20510 standard; protein; 1998 AA.

XX AAE20510;

DT 01-JUL-2002 (first entry)

DE Human ion channel protein #1.

XX Human; novel human protein; NHP; voltage-gated sodium channel;
KW gene therapy; bioreactor; mental disorder; biological disorder;
KW medical disorder.

XX Homo sapiens.

OS

XX Key Location/Qualifiers

FT Misc-difference 981 /note= "Encoded by MTG"

FT Misc-difference 1056 /note= "Encoded by RCA"

XX WO200214498-A2.

PN 21-FEB-2002.

PF 15-AUG-2001; 2001WO-US025650.

PR 16-AUG-2000; 2000US-0225989P.

PA (LEXI-) LEXICON GENETICS INC.

PI Turner CA, Mathur B, Mathur D;

XX WPI; 2002-280757/32.

DR N-PSDB; AAD32839.

PT Novel polynucleotides encoding human sodium channel proteins,
particularly voltage-gated sodium channel proteins useful for drug
screening, diagnosis and in gene therapy of biological disorders.
PS Claim 1; Page 30-34; 83pp; English.

CC The present sequence is novel human protein (NHP), ion channel protein.
CC NHP share structural similarity with mammalian sodium channel proteins
CC particularly voltage-gated sodium channel proteins. NHP oligonucleotides
CC are useful as hybridisation probes for screening libraries and assessing
CC gene expression patterns. Sequences derived from regions adjacent to the

CC intron/exon boundaries of NHP gene can be used to design primers for use
 CC in amplification assays to detect mutations within the exons, splice
 CC sites, introns that can be used in diagnostics and pharmacogenomics. NHP
 CC nucleotide sequences are useful for drug screening effective in the
 CC treatment of symptomatic or phenotypic manifestations of perturbing the
 CC normal function of NHP in the body, and nucleotide constructs encoding the
 CC NHP products are useful to genetically engineer host cells to express NHP
 CC products in vivo. These genetically engineered cells function as
 CC bioreactors in the body delivering a continuous supply of a NHP, a NHP
 CC peptide, or a NHP fusion protein to the body. Nucleotide construct
 CC encoding NHP products are also useful in gene therapy for modulating NHP
 CC expression and to produce genetically engineered host cells to express
 CC NHP products in vivo. NHP nucleotide sequences may also be used as part
 CC of ribozyme and/or triple helix sequences that are useful for NHP gene
 CC regulation. The NHP polypeptides are useful for generating antibodies, as
 CC reagents in diagnostic assays, for identifying other cellular gene
 CC products related to NHP and as reagents in assays for screening for
 CC compounds that are useful in the treatment of mental, biological or
 CC medical disorders and diseases

CC Sequence 1998 AA;
 SQ

Query Match 100.0%; Score 28; DB 5; Length 1998;
 Best Local Similarity 100.0%; Pred. No. 1.4e-20;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 MYLVEVFIIIFGSPFTLNLFIGVIIDNF 28
 DB 1448 MYLVEVFIIIFGSPFTLNLFIGVIIDNF 1475

RESULT 17
 ABR83184
 ID ABR83184 standard; protein; 1998 AA.
 AC ABR83184;
 XX

DT 15-JAN-2004 (first entry)
 XX

DE Human SCN1A splice variant -33P:SCN1ADELP671-681.
 XX

KM SCN1A; sodium channel type 1 alpha-subunit; anticonvulsant; analgesic;
 KM neuroprotective; anesthetic; cyostatic; cerebroprotective; cardiac;
 KM hypotensive; gene therapy; human; splice variant.
 OS Homo sapiens.

XX MO2003072751-A2.
 XX

PD 04-SEP-2003.
 XX

PF 25-FEB-2003; 2003WO-US006010.
 XX

PR 25-FEB-2002; 2002US-0359382P.
 XX

PA (UYVA-) UNIV VANDERBILT.
 XX

PI George AL, Lossin C;
 XX

DR WPI; 2003-712725/67.
 XX

DR N-PSDB; ACF57879.
 XX

PT Recombinantly expressed sodium channel type 1 alpha subunit, useful in
 screening for modulators, for treating e.g. epilepsy.
 XX

PS Disclosure; Page 148-156; 176pp; English.
 XX

CC The invention relates to a recombinantly expressed and isolated human
 CC SCN1A (sodium channel type 1 alpha-subunit) (I). (I), optionally
 CC incorporated into a cell, is used to screen for specific modulators,
 CC potentially useful as anticonvulsant, antiepileptic, neuroprotective,
 CC analgesic and/or anesthetic agents, e.g. for treating severe myoclonic
 CC epilepsy of infancy, stroke, cardiac arrest, hyperkalemic paralysis,

CC motor endplate diseases, hypertension, congestive heart failure and
 CC muscular dystrophy also to treat cancer (SCN1A is expressed in prostatic
 CC and metastatic cancer cell lines). These activities can also be provided
 CC by gene therapy vectors that express (I) or the modulators. The
 CC modulators, also antibodies directed against (I), are used to detect
 CC sodium channel polypeptides. The present sequence represents a human
 CC SCN1A splice variant 33P:SCN1ADELP671-681 encoding cDNA
 XX
 SQ Sequence 1998 AA;

Query Match 100.0%; Score 28; DB 7; Length 1998;
 Best Local Similarity 100.0%; Pred. No. 1.4e-20;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 MYLVEVFIIIFGSPFTLNLFIGVIIDNF 28
 DB 1448 MYLVEVFIIIFGSPFTLNLFIGVIIDNF 1475

RESULT 18
 ABB06026

ID ABB06026 standard; protein; 1999 AA.
 AC ABB06026;
 XX

XX 10-MAY-2002 (first entry)
 DT

DE Human sodium channel SCN1A protein SEQ ID NO:2.
 XX

KM Human; sodium channel; SCN1A; chromosome 2q24;
 KM familial hypocalcaemic periodic paralysis; motor endplate disease.
 OS Homo sapiens.

XX MO200196552-A1.
 XX

PD 20-DEC-2001.
 XX

PF 12-JUN-2001; 2001WO-JP004956.
 XX

PR 13-JUN-2000; 2000JP-00177540.
 XX

PR 13-JUN-2000; 2000JP-00177544.
 XX

PA (NISC-) JAPAN SCI & TECHNOLOGY CORP.
 XX

PI Kanazawa I, Goto J, Jeong S;
 XX

DR WPI; 2002-098066/13.
 XX

DR N-PSDB; ABL39689.
 XX

PT Human sodium channels SCN1A and SCN3A and encoded genes, useful in
 studying physiological mechanism in which excitant cells participate and
 causes of diseases and developing drugs for motor endplate disease.
 XX
 PS Claim 1; Page 40-49; 88pp; Japanese.
 XX

CC The present invention describes human sodium channels SCN1A and SCN3A.
 CC The present sequence represents the human sodium channel SCN1A. SCN1A and
 CC SCN3A have been located to the human chromosome 2 long arm, positions
 CC 2q24 and 2q24-31 respectively. The sodium channel proteins are useful in
 CC studying the physiological mechanism in which excitant cells participate
 CC and cause human diseases, and in developing remedies for e.g. familial
 CC hypocalcaemic periodic paralysis of extremities and motor endplate
 CC disease
 XX

SQ Sequence 1999 AA;

Query Match 100.0%; Score 28; DB 5; Length 1999;
 Best Local Similarity 100.0%; Pred. No. 1.4e-20;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 MYLVEVFIIIFGSPFTLNLFIGVIIDNF 28
 ||||||||||||||||||||||||||||

XX Rouleau GA, Lafreniere RG, Rochefort D, Cossette P, Ragsdale D;
 PI WPI; 2001-355945/37.
 DR N-PSDB; AAH55793.
 XX
 PT Determining a predisposition to epilepsy and/or development of epilepsy
 PT comprises determining the genotype of SCN1A, SCN2A and/or SCN3A, or a DNA
 PT variant, equivalent, or mutation which shows a linkage disequilibrium.
 PS Disclosure; Page 123-130; 268pp; English.

XX The present invention describes a method (M1) of determining an
 CC individual's predisposition to epilepsy and/or development of epilepsy,
 CC as well as predicting the individual's response to medication. The method
 CC comprises determining the genotype of at least one gene selected from
 CC SCN1A, SCN2A or SCN3A, or a DNA variant, equivalent, or mutation which
 CC shows a linkage disequilibrium. SCN1A, SCN2A and SCN3A are all sodium
 CC channel genes located on chromosome 2. The idiopathic generalised
 CC epilepsy (IGE) gene is more specifically localised on chromosome 2q23-
 CC q31. Compounds identified as modulators of the biological activity of
 CC SCN1A, SCN2A or SCN3A proteins or genes, are useful for treating epilepsy
 CC or other neurological disorders. They have anticonvulsant and
 CC neuroprotective activities. AAH55763 to AAH56164 and AAB9674 to AAB9679
 CC represent SCN1A, SCN2A, and SCN3A cDNAs, gene fragments, PCR primers,
 CC oligonucleotides and proteins given in the exemplification of the present
 CC invention

SQ Sequence 2005 AA;

Query Match 100.0%; Score 28; DB 4; Length 2005;
 Best Local Similarity 100.0%; Pred. No. 1.4e-20;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 MYLYEVFIIFGSPFTLNLFIVGIIDNF 28
 |||||
 DB 1449 MYLYEVFIIFGSPFTLNLFIVGIIDNF 1476

RESULT 22
 AAB9677 standard; protein; 2005 AA.

AC AAB9677;
 XX
 DT 04-SEP-2001 (first entry)
 XX

DE Human neonatal form of SCN2A protein sequence SEQ ID NO:36.

XX Human; epilepsy; chromosome 2; SCN1A; SCN2A; SCN3A; identification;
 KW diagnosis; mutation; chromosome 2q23-q31; neurological disorder;
 KW anticonvulsant; neuroprotective.

OS Homo sapiens.
 XX
 PN WO200138564-A2.
 XX

PD 31-MAY-2001.

PF 24-NOV-2000; 2000WO-CA001404.

PR 26-NOV-1999; 99US-0167623P.

PA (UYMC-) UNIV MCGILL.

PI Rouleau GA, Lafreniere RG, Rochefort D, Cossette P, Ragsdale D;
 XX WPI; 2001-355945/37.
 DR N-PSDB; AAH55794.
 XX

PT Determining a predisposition to epilepsy and/or development of epilepsy
 PT comprises determining the genotype of SCN1A, SCN2A and/or SCN3A, or a DNA
 PT variant, equivalent, or mutation which shows a linkage disequilibrium.

XX Disclosure; Page 131-138; 268pp; English.

XX The present invention describes a method (M1) of determining an
 CC individual's predisposition to epilepsy and/or development of epilepsy,
 CC as well as predicting the individual's response to medication. The method
 CC comprises determining the genotype of at least one gene selected from
 CC SCN1A, SCN2A or SCN3A, or a DNA variant, equivalent, or mutation which
 CC shows a linkage disequilibrium. SCN1A, SCN2A and SCN3A are all sodium
 CC channel genes located on chromosome 2. The idiopathic generalised
 CC epilepsy (IGE) gene is more specifically localised on chromosome 2q23-
 CC q31. Compounds identified as modulators of the biological activity of
 CC SCN1A, SCN2A or SCN3A proteins or genes, are useful for treating epilepsy
 CC or other neurological disorders. They have anticonvulsant and
 CC neuroprotective activities. AAH55763 to AAH56164 and AAB9674 to AAB9679
 CC represent SCN1A, SCN2A, and SCN3A cDNAs, gene fragments, PCR primers,
 CC oligonucleotides and proteins given in the exemplification of the present
 CC invention

SQ Sequence 2005 AA;

Query Match 100.0%; Score 28; DB 4; Length 2005;
 Best Local Similarity 100.0%; Pred. No. 1.4e-20;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 MYLYEVFIIFGSPFTLNLFIVGIIDNF 28
 |||||
 DB 1449 MYLYEVFIIFGSPFTLNLFIVGIIDNF 1476

RESULT 23
 ABB83627 standard; protein; 2005 AA.

AC ABB83627;

DT 10-OCT-2002 (first entry)

DE Human GERS+ protein with SCN2A mutation.

XX Human; GERS+; SCN2A; mutant; mutein;
 KW generalized epilepsy with febrile seizure plus.

OS Homo sapiens.

PN JP2002136289-A.

PD 14-MAY-2002.

PF 01-NOV-2000; 2000JP-00334969.

PR 01-NOV-2000; 2000JP-00334969.

PA (KAGA-) KAGAKU GIYUSU SHINKO JIGYODAN.
 PA (RIKA) RIKAGAKU KENKYUSHO.

PI WPI; 2002-552308/59.
 DR N-PSDB; ABQ79201.

PT A human polynucleotide which is complementary to an mRNA transcribed from
 PT a generalized epilepsy with febrile seizure plus (GERS+)-related gene
 PT useful for diagnosing GERS+.

PS Claim 10; Page 29-34; 37pp; Japanese.

XX This invention relates to a human polynucleotide which is complementary
 CC to an mRNA transcribed from a "generalized epilepsy with febrile seizure
 CC plus" (GERS+)-related gene. The gene is useful for diagnosing GERS+. The
 CC present sequence represents the human GERS+ protein sequence with SCN2A
 CC mutation

SQ Sequence 2005 AA;

Query Match 100.0%; Score 28; DB 5; Length 2005;
 Best Local Similarity 100.0%; Pred. No. 1,4e-20;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 MYLYFVIRIIFGSPFTLNLFIGVIIDNF 28
 |||||
 1449 MYLYFVIRIIFGSPFTLNLFIGVIIDNF 1476

RESULT 24
 ADB78604
 ID ADB78604 standard; protein; 2005 AA.

XX ADB78604;

DT 04-DEC-2003 (first entry)

XX Human sodium channel subunit mutant SEQ ID NO:148.

KW murein; mutant; ion channel; ion channel subunit; ICS; nootropic;
 KW neuroprotective; inotropic; antipyretic; antiarrhythmic; antimigraine;
 KW antidepressant; antiparkinsonian; neuroleptic; tranquilizer; analgesic;
 KW nephrotoxic; antidiabetic; ophthalmological; epilepsy;
 KW ion channel dysfunction; human.

XX Synthetic.
 OS Homo sapiens.

XX WO2003008574-A1.

XX 30-JAN-2003.

XX 08-JUL-2002; 2002WO-AU000910.

XX 18-JUL-2001; 2001AU-00006452.

XX 05-MAR-2002; 2002AU-00000910.

XX 13-MAY-2002; 2002AU-00002292.

XX (BION-) BIONOMICS LTD.

XX (WALL/) WALLACE R W.

XX Mulley JC, Harkin LA, Dibbens LM, Phillips HA, Heron SE;

XX Berkovic SF, Scheffer IE;

XX WPI; 2003-239332/23.

XX N-PSDB; ADB78643.

XX Claim 13; SEQ ID NO 148; 106pp; English.

The invention relates to a novel method for identifying a subject predisposed to a disorder associated with ion channel dysfunction. The method comprises ascertaining if at least one of the genes encoding ion channel subunits (ICS) has undergone a mutation event so that a cDNA derived from the subject has any of 134 nucleotide sequences. The method of the invention has nootropic, neuroprotective, inotropic, antipyretic, antiarrhythmic, antimigraine, antidepressant, antiparkinsonian, neuroleptic, tranquilizer, analgesic, nephrotoxic, antidiabetic, and ophthalmological activity. A polynucleotide of the invention acts as an ion channel agonist, or ion channel antagonist. The methods, isolated nucleic acids, polypeptides, antibody, selective agonist, antagonist or modulator of an ion channel, cells and genetically modified non-human animal, are useful for the diagnosis and treatment of epilepsy and/or a disorder associated with ion channel dysfunction, such as hyper- or hypokalemic periodic paralysis, myotonia, malignant hyperthermia, CC myasthenia, cardiac arrhythmias, episodic ataxia, migraine, Alzheimer's disease, Parkinson's disease, schizophrenia, hyperkplexia, anxiety, depression, phobic obsessive symptoms, neuropathic pain, inflammatory pain, chronic/acute pain, Barter's syndrome, polycystic kidney disease,

CC Dent's disease, hyperinsulinaemic hypoglycaemia of infancy, cystic
 CC fibrosis, congenital stationary night blindness and total colour
 CC blindness. The present sequence represents a mutant protein of the
 CC invention. The sequence data for this patent is not represented in the
 CC printed specification, but was obtained in electronic format directly
 CC from WIPO at ftp.wipo.int/pub/published_pat_sequences.

XX Sequence 2005 AA;

Query Match 100.0%; Score 28; DB 7; Length 2005;
 Best Local Similarity 100.0%; Pred. No. 1,4e-20;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 MYLYFVIRIIFGSPFTLNLFIGVIIDNF 28
 |||||
 Db 1449 MYLYFVIRIIFGSPFTLNLFIGVIIDNF 1476

RESULT 25

ADB78603
 ID ADB78603 standard; protein; 2005 AA.

XX ADB78603;

DT 04-DEC-2003 (first entry)

XX Human sodium channel subunit mutant SEQ ID NO:147.

KW murein; mutant; ion channel; ion channel subunit; ICS; nootropic;
 KW neuroprotective; inotropic; antipyretic; antiarrhythmic; antimigraine;
 KW antidepressant; antiparkinsonian; neuroleptic; tranquilizer; analgesic;
 KW nephrotoxic; antidiabetic; ophthalmological; epilepsy;
 KW ion channel dysfunction; human.

XX Synthetic.

XX Homo sapiens.

XX WO2003008574-A1.

XX 30-JAN-2003.

XX 08-JUL-2002; 2002WO-AU000910.

XX 18-JUL-2001; 2001AU-00006452.

XX 05-MAR-2002; 2002AU-00000910.

XX 13-MAY-2002; 2002AU-00002292.

XX (BION-) BIONOMICS LTD.

XX (WALL/) WALLACE R W.

XX Mulley JC, Harkin LA, Dibbens LM, Phillips HA, Heron SE;

XX Berkovic SF, Scheffer IE;

XX WPI; 2003-239332/23.

XX N-PSDB; ADB78642.

XX Claim 13; SEQ ID NO 147; 106pp; English.

The invention relates to a novel method for identifying a subject predisposed to a disorder associated with ion channel dysfunction. The method comprises ascertaining if at least one of the genes encoding ion channel subunits (ICS) has undergone a mutation event so that a cDNA derived from the subject has any of 134 nucleotide sequences. The method of the invention has nootropic, neuroprotective, inotropic, antipyretic, antiarrhythmic, antimigraine, antidepressant, antiparkinsonian, neuroleptic, tranquilizer, analgesic, nephrotoxic, antidiabetic, and ophthalmological activity. A polynucleotide of the invention acts as an ion channel agonist, or ion channel antagonist. The methods, isolated

CC nucleic acids, polypeptides, antibody, selective agonist, antagonist or
 CC modulator of an ion channel, cells and genetically modified non-human
 CC animal, are useful for the diagnosis and treatment of epilepsy and/or a
 CC disorder associated with ion channel dysfunction, such as hyper- or hypo-
 CC kalemic periodic paralysis, myotonia, malignant hyperthermia,
 CC myasthenia, cardiac arrhythmias, episodic ataxia, migraine, Alzheimer's
 CC disease, Parkinson's disease, schizophrenia, hyperkplexia, anxiety,
 CC depression, phobic obsessive symptoms, neuropathic pain, inflammatory
 CC pain, chronic/acute pain, Bartter's syndrome, polycystic kidney disease,
 CC Dent's disease, hyperinsulinaemic hypoglycaemia of infancy, cystic
 CC fibrosis, congenital stationary night blindness and total colour
 CC blindness. The present sequence represents a mutant protein of the
 CC invention. The sequence data for this patent is not represented in the
 CC printed specification, but was obtained in electronic format directly
 CC from WIPO at ftp.wipo.int/pub/published_pat_sequences.
 XX

Sequence 2005 AA;

Query Match 100.0%; Score 28; DB 7; Length 2005;
 Best Local Similarity 100.0%; Pred. No. 1.4e-20;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 MYLYFVIFIIIPGSPFTLNLFIGVITDNE 28
 |||||
 Db 1449 MYLYFVIFIIIPGSPFTLNLFIGVITDNE 1476

Search completed: January 27, 2005, 17:45:16
 Job time : 87.5 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 27, 2005, 17:36:05 ; Search time 17 Seconds
(without alignments)
158.475 Million cell updates/sec

Title: US-10-608-584-19

Perfect score: 28
Sequence: 1 MYLYVFIFIFGSPFTLNLFGVINDF 28

Scoring table: OLIGO
Gapop 60.0 , Gapext 60.0

Searched: 283416 seqs, 96216763 residues

Word size: 0

Total number of hits satisfying chosen parameters: 283416

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Listing first 100 summaries

Database: PIR_79.*

1: p1r1.*
2: p1r2.*
3: p1r3.*
4: p1r4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	28	100.0	1835	2	154333 sodium channel alp
2	28	100.0	1836	2	164893 sodium channel alp
3	28	100.0	1836	2	JS0648 sodium channel alp
4	28	100.0	1836	2	IS1964 sodium channel alp
5	28	100.0	1840	1	CHRTM1 sodium channel pro
6	28	100.0	1951	2	S00320 sodium channel pro
7	28	100.0	1983	2	A60054 sodium channel pro
8	28	100.0	2005	2	A46269 sodium channel alp
9	28	100.0	2005	2	B25019 sodium channel pro
10	28	100.0	2009	2	A25019 sodium channel pro
11	25	89.3	1976	2	IS6555 sodium channel pro
12	25	89.3	2016	2	A38195 sodium channel pro
13	21	75.0	2295	2	A56595 sodium channel alp
14	21	75.0	1689	2	S72467 voltage-gated sodi
15	21	75.0	1699	2	T31340 sodium channel alp
16	21	75.0	1765	2	T42388 sodium channel alp
17	21	75.0	1784	2	T43167 sodium channel pro
18	21	75.0	1820	2	A33299 sodium channel pro
19	21	75.0	1977	2	S54771 sodium channel alp
20	21	75.0	2019	2	A33996 sodium channel pro
21	21	75.0	2108	2	S72458 sodium channel pro
22	18	64.3	2049	2	T43161 sodium channel pro
23	17	60.7	1321	2	A60165 sodium channel pro
24	17	60.7	1810	2	T31092 probable voltage-g
25	15	53.6	1820	1	CHER sodium channel pro
26	13	46.4	1993	2	T30902 sodium channel SCA
27	12	42.9	1695	2	JB0084 voltage-gated sodi
28	12	42.9	1739	2	A48298 sodium channel hom
29	8	28.6	1522	2	JC1101 sodium channel pro

30	7	25.0	260	2	G81290 probable capsule p
31	7	25.0	274	2	B83837 hypothetical prote
32	7	25.0	351	2	A69808 H+/Ca2+ exchanger
33	7	25.0	399	2	B84964 hypothetical prote
34	7	25.0	491	2	T23524 hypothetical prote
35	7	25.0	494	1	O4H0A6 coumarin-7-hydroxy
36	7	25.0	542	2	F90457 hypothetical prote
37	7	25.0	568453	2	sodium channel pro
38	6	1957	1957	2	legumain glycoprot
39	6	21.4	22	2	S17303 60S ribosomal prot
40	6	21.4	118	2	C90105 hypothetical prote
41	6	21.4	129	2	T25141 conserved hypothet
42	6	21.4	130	2	C90503 probable membrane
43	6	21.4	153	2	S52605 hypothetical prote
44	6	21.4	155	2	A91055 hypothetical prote
45	6	21.4	155	2	D85899 hypothetical prote
46	6	21.4	162	2	T13574 Mdh2 dehydrogenas
47	6	21.4	164	2	F65031 hypothetical prote
48	6	21.4	182	1	R5MX6 ribosomal protein
49	6	21.4	182	2	A45553 22K surface membra
50	6	21.4	184	2	B90592 hypothetical prote
51	6	21.4	192	2	S09506 hypothetical prote
52	6	21.4	218	2	S43591 hypothetical prote
53	6	21.4	220	2	B64681 nicotinamide monon
54	6	21.4	220	2	B71832 hypothetical prote
55	6	21.4	221	2	D86878 hypothetical prote
56	6	21.4	237	2	S64315 probable replicati
57	6	21.4	240	2	B84497 H+-transporting tw
58	6	21.4	248	2	S57461 hypothetical prote
59	6	21.4	251	2	S38811 signal peptidase t
60	6	21.4	254	2	B97242 probable ABC trans
61	6	21.4	256	2	AG0413 flagellar biosynth
62	6	21.4	258	2	B72319 probable permease
63	6	21.4	259	2	A83294 hypothetical prote
64	6	21.4	261	2	AD1721 hypothetical prote
65	6	21.4	261	2	F90506 hypothetical prote
66	6	21.4	279	2	B70332 hypothetical prote
67	6	21.4	282	2	C84114 UDP-glucose 4-epim
68	6	21.4	286	2	I46855 voltage-gated pota
69	6	21.4	314	2	T32672 hypothetical prote
70	6	21.4	316	2	S46055 probable membrane
71	6	21.4	317	2	A71698 hypothetical prote
72	6	21.4	322	2	B81279 enterochelin uptak
73	6	21.4	322	2	D96921 oligopeptide trans
74	6	21.4	331	1	O4H0B8 cytochrome P450 2A
75	6	21.4	331	2	B84195 hypothetical prote
76	6	21.4	335	2	G83598 cell division prot
77	6	21.4	336	2	F75320 WD-repeat family p
78	6	21.4	339	2	S61020 probable membrane
79	6	21.4	340	2	S41752 UDP-3-O-(3-hydroxy
80	6	21.4	340	2	AE0129 UDP-3-O-(3-hydroxy
81	6	21.4	341	2	AD0530 UDP-3-O-(3-hydroxy
82	6	21.4	341	2	S13729 UDP-3-O-(3-hydroxy
83	6	21.4	341	2	B85502 hypothetical prote
84	6	21.4	341	2	E90651 hypothetical prote
85	6	21.4	341	2	B37083 Ssc protein - Salm
86	6	21.4	345	2	T12354 Mdh2 dehydrogenas
87	6	21.4	351	2	AF3768 microcin C7 self-i
88	6	21.4	365	2	AF3119 hypothetical prote
89	6	21.4	374	2	A98168 hypothetical prote
90	6	21.4	375	2	D82894 heat shock protein
91	6	21.4	381	2	S59093 ubiquinol-cytochro
92	6	21.4	398	2	D75093 macroide-efflux d
93	6	21.4	406	2	T23934 hypothetical prote
94	6	21.4	409	2	AG0235 probable sugar tra
95	6	21.4	409	2	T47026 hypothetical prote
96	6	21.4	411	2	B84949 tetrahydrofolylpol
97	6	21.4	434	2	AH1990 hypothetical prote
98	6	21.4	437	2	G82777 glutamate symport
99	6	21.4	444	2	F71916 probable lipopolys
100	6	21.4	468	2	S75389 probable phenylala

channel alpha subunit - hmsc

C/Species: Homo sapiens (man)
C/Date: 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change 20-Aug-1999
C/Accession: I54I23
R/MC/Catchey, A.I.; Lin, C.S.; Wang, J.; Hoffman, E.P.; Rojas, C.; Gusella, J.F.
Hum. Mol. Genet. 1: 521-527, 1992
A>Title: The genomic structure of the human skeletal muscle sodium channel gene.
A/Accession: I54I23; NCID:93338444; PMID:1339144
A>Status: preliminary; translated from GB/EMBL/DBJ
A/Molecule type: DNA
A/Residues: 1-1835 <RES>
A/Cross-references: GB:L01983; NID:G337992; PIDN:AA75557.1; PID:9908809
A/Genetics:
A/Genes: GDB:SCNA4
A/Cross-references: GDB:125181; OMIM:170500
A/Map position: 17q23.1-17q25.3
A/Introns: 91/3; 131/2; 161/2; 204/2; 235/1; 346/1; 367/2; 414/3; 484/3; 536/1; 615/3; 616/3
A/Superfamily: sodium channel protein
C/Keywords: duplication

Matches 28; Conservatively 0; Pred. No. 3.4e-20;

Qy	1	MYLVYVIFIFIGSFFTLNLFGVIIND	28	0	0	0
Db	1271	MYLVYVIFIFIGSFFTLNLFGVIIND	1298			

sodium channel alpha subunit - human

C:Species: Homo sapiens (man)
C:Date: 06-Sep-1996 #sequence_revision 06-Sep-1996 #text_change 09-Jul-2004
C:Accession: I64893
R:George, A.L.
Ann: Neurol. 31, 131-137, 1992
A:Title: Primary structure of the adult human skeletal muscle voltage-dependent sodium c
A:Reference number: I51964; MUID:92246457; PMID:1315496
A:Accession: I64893
A:Status: Preliminary; translated from GB/EMBL/DBJ.
A:Molecule type: mRNA
A:Residues: 1-1836 <RES>
A:Cross-references: UNIPROT:P35499; GB:M81758; NID:G338212; PIDN:AAA60554.1; PID:G338213
C:Genetics
A:Gene: SKM1
C:Superfamily: sodium channel protein
C:Keywords: duplication

Matches 28; Conservatively 0.0%; Pred. No. 3.4e-20;

Qy	1	MYLYEVIFIIFGSFFTLNFIGVIIDNF	28	Indels	Gaps
Dy	1271	MYLYEVIFIIFGSFFTLNFIGVIIDNF	1298		

sodium channel alpha chain - human

C:Species: Homo sapiens (man) human
C:Date: 30-Jun-1992 #sequence
C:Accession: J06648-A4209
R:Wang, J.; Rojars, C.V.; Zhou, J.; Schwartz, L.S.; Nicholas, H.; Hoffmann, E.P.
Biochem. Biophys. Res. Commun. 182, 794-801, 1992
A:Title: Sequence and genomic structure of the human adult skeletal muscle sodium channel

A:Reference number: US0648; MUID:92134303; PMID:1310396
A:Accession: JS0648
A:Status: nucleic acid sequence not shown
A:Molecule type: mRNA
A:Residues: 1-1836 <MAN>
A:Cross-references: UNIPROT:P35499
A:Note: 861-Aap was also found as the result of polymorphism
R:McClatchey, A. I.; Van den Bergh, P.; Pericak-Vance, M. A.; Raskind, W.; Vervellen, C.; Mc
Cell 68, 769-774, 1992
A:Title: Temperature-sensitive mutations in the III-IV cytoplasmic loop region of the s
A:Reference number: A42099; MUID:92154589; PMID:1310898
A:Accession: A42099
A:Molecule type: DNA
A:Residues: 1299-1351 <MCC>
A:Cross-references: GB:S82622; NID:9245611; PIDN:AAB21450.1; PID:9245612
A:Experimental source: skeletal muscle
A:Note: sequence extracted from NCBI backbone (NCBIN:82622, NCBIID:82623)
Identics:

```

References: GDB:125181; OMIM:170500
A;Map position: 17q23.1-17q25.3

```

C:Superfamily: sodium channel protein
C:Keywords: duplication, glycoprotein, phosphoprotein, transmembrane protein
F:129-150/Domain: transmembrane #status predicted <R1>
F:159-178/Domain: transmembrane #status predicted <R2>
F:191-210/Domain: transmembrane #status predicted <R3>
F:217-236/Domain: transmembrane #status predicted <R4>
F:253-266/Domain: transmembrane #status predicted <R5>
F:424-449/Domain: transmembrane #status predicted <R6>
F:574-597/Domain: transmembrane #status predicted <R7>
F:609-632/Domain: transmembrane #status predicted <R8>
F:641-660/Domain: transmembrane #status predicted <R12>
F:667-686/Domain: transmembrane #status predicted <R13>
F:702-724/Domain: transmembrane #status predicted <R14>
F:777-802/Domain: transmembrane #status predicted <R15>
F:1027-1049/Domain: transmembrane #status predicted <R16>
F:1064-1089/Domain: transmembrane #status predicted <R11>
F:1096-1116/Domain: transmembrane #status predicted <R12>
F:1122-1143/Domain: transmembrane #status predicted <R13>
F:1163-1184/Domain: transmembrane #status predicted <R14>
F:1269-1295/Domain: transmembrane #status predicted <R15>
F:1349-1372/Domain: transmembrane #status predicted <R16>
F:1384-1407/Domain: transmembrane #status predicted <R11>
F:1414-1437/Domain: transmembrane #status predicted <R12>
F:1447-1469/Domain: transmembrane #status predicted <R13>
F:1485-1507/Domain: transmembrane #status predicted <R14>
F:1574-1598/Domain: transmembrane #status predicted <R15>
F:1621-1649,220,378,415,1019,1130,1242,1313,1721,1826/Binding site: phosphate (Thr) (covalent) (by protein kinase A) #status predicted <V86>
F:156,251,553,553,553,553,1571,1146/Binding site: phosphate (Ser) (covalent) (by protein kinase A) #status predicted <V87>
F:2121,288,291,297,303,315,321,333,366,507,702,961,1191,1205/Binding site: phosphate (Ser) (covalent) (by protein kinase A) #status predicted <V88>
F:3246,670,725,850,950,1127,1195,1338/Binding site: phosphate (Ser) (covalent) (by protein kinase A) #status predicted <V89>
F:367,457/Binding site: phosphate (Thr) (covalent) (by protein kinase A) #status predicted <V90>

```

- - conservative; mismatches 0; indels

```

QY 1 MYLYFVFIIFGSEFTLNLFIGVIIDNF 28
 |||||
 Db 1271 MYLYFVFIIFGSEFTLNLFIGVIIDNF 1298
 |||||

sodium channel alpha chain SCN4A, skeletal muscle

Ci:Species: Homo sapiens (man)
Ci>Date: 24-May-1996 #sequence_revision 24-May-1996 #text_change 09-Jul-2004
Ci:Accession: IS1964
R:George, A.L.
Ann. Neurol. 31, 131-137, 1992
A/Title: Primary structure of the adult human skeletal muscle voltage-dependent sodium ch
A/Reference number: IS1964, PMID:92246457, PMID:1515496
A/Accession: IS1964

A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-1836 <RES>
A:Cross-references: UNIPROT:P35499; GB:L04236; NID:G9292485; PIDN:AAB59624.1; PID:G9292487
C:Genetics:
A:Gene: GDB:SCN4A
A:Cross-references: GDB:125181; OMIM:170500
A:Map position: 17q23.1-17q25.3
A:Insertions: 91/3; 131/2; 161/2; 204/2; 235/1; 346/1; 367/2; 414/3; 484/3; 536/1; 615/3; 6
C:Superfamily: sodium channel protein
C:Keywords: duplication; skeletal muscle

Query Match 100.0%; Score 28; DB 2; Length 1836;
Best Local Similarity 100.0%; Pred. No. 3.4e-20;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 MYLYFVFIIFGSPFTLNLFGVIIDNF 28
Db 1271 MYLYFVFIIFGSPFTLNLFGVIIDNF 1298

RESULT 5
CHR1M1
sodium channel protein mu1 alpha chain, skeletal muscle - rat
C:Species: Rattus norvegicus (Norway rat)
C>Date: 30-Sep-1990 #sequence_revision 30-Sep-1990 #text_change 09-Jul-2004
C:Accession: JN0007
R:Trimmer, J.S.; Cooperman, S.S.; Tomiko, S.A.; Zhou, J.; Crean, S.M.; Boyle, M.B.; Kall
Neuron 3, 33-49, 1989
A:Title: Primary structure and functional expression of a mammalian skeletal muscle sodi
A:Reference number: JN0007; MUID:90148778; PMID:2559760
A:Accession: JN0007
A:Molecule type: mRNA
A:Residues: 1-1840 <TRI>
A:Cross-references: UNIPROT:P15390; GB:M26643; NID:G205651; PIDN:AAA41682.1; PID:G205652
C:Comment: Action potentials propagated by skeletal muscle sodium channels are responsib
C:Comment: This heavily glycosylated protein contains four homologous domains, each of w
C:Comment: This protein is distinct from but related to sodium channel proteins isolated
C:Superfamily: sodium channel protein
C:Keywords: duplication; glycoprotein; ion transport; neuromuscular junction; phosphoprc
F:120-458,561-813,1013-1305,1335-1611,Region: duplication
F:56,251,1321,1504/Binding site: phosphatase (Ser) (covalent) (by CAMP-dependent kinase) #
F:214,288,291,297,303,309,315,327,356,502,696,954,1184,1198,1563,1702/Binding site: catr

Query Match 100.0%; Score 28; DB 1; Length 1840;
Best Local Similarity 100.0%; Pred. No. 3.4e-20;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 MYLYFVFIIFGSPFTLNLFGVIIDNF 28
Db 1264 MYLYFVFIIFGSPFTLNLFGVIIDNF 1291

RESULT 6
S00320
sodium channel protein III - rat
C:Species: Rattus norvegicus (Norway rat)
C>Date: 30-Jun-1989 #sequence_revision 30-Jun-1989 #text_change 09-Jul-2004
C:Accession: S00320
R:Kayano, T.; Noda, M.; Flockerzi, V.; Takahashi, H.; Numa, S.
FEBS Lett. 228, 187-194, 1988
A:Title: Primary structure of rat brain sodium channel III deduced from the cDNA sequenc
A:Reference number: S00320; MUID:88137594; PMID:2449363
A:Accession: S00320
A:Molecule type: mRNA
A:Residues: 1-1951 <RAY>
A:Cross-references: UNIPROT:P08104; EMBL:Y00766; NID:G57210; PIDN:CA68735.1; PID:G57211
A:Note: 270-Ile, 278-Ileu, 355-Lys, and 1055-Arg were also found
C:Superfamily: sodium channel protein
C:Keywords: duplication; transmembrane protein

Query Match 100.0%; Score 28; DB 2; Length 1951;
Best Local Similarity 100.0%; Pred. No. 3.6e-20;

Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 MYLYFVFIIFGSPFTLNLFGVIIDNF 28
Db 1395 MYLYFVFIIFGSPFTLNLFGVIIDNF 1422

RESULT 7
A60054
sodium channel protein IIB, long form - rat
C:Species: Rattus norvegicus (Norway rat)
C>Date: 03-Mar-1993 #sequence_revision 03-Mar-1993 #text_change 09-Jul-2004
C:Accession: A60054; B44824
R:Joho, R.H.; Moorman, J.R.; Vandongen, A.M.J.; Kirsch, G.E.; Silberberg, H.; Schuster,
Brain Res. Mol. Brain Res. 7, 105-113, 1990
A:Title: Toxin and kinetic profile of rat brain type IIB sodium channels expressed in Xe
A:Reference number: A60054; MUID:90251117; PMID:2160038
A:Accession: A60054
A:Status: not compared with conceptual translation
A:Molecule type: mRNA
A:Residues: 1-1983 <JOH>
A:Cross-references: UNIPROT:Q64243
R:Schaller, K.U.; Krzemien, D.M.; McKenna, N.M.; Caldwell, J.H.
J. Neurosci. 12, 1370-1381, 1992
A:Title: Alternatively spliced sodium channel transcripts in brain and muscle.
A:Reference number: A44824; MUID:92211397; PMID:1313493
A:Accession: B44824
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 611-662 <SCH>
A:Cross-references: GB:S97388; NID:G248225; PIDN:AAB21984.1; PID:G248226
A:Experimental source: skeletal muscle
A:Note: sequence inconsistent with the nucleotide translation
C:Superfamily: sodium channel protein
C:Keywords: duplication; glycoprotein; ion transport; sodium channel; transmembrane prot

Query Match 100.0%; Score 28; DB 2; Length 1983;
Best Local Similarity 100.0%; Pred. No. 3.6e-20;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 MYLYFVFIIFGSPFTLNLFGVIIDNF 28
Db 1427 MYLYFVFIIFGSPFTLNLFGVIIDNF 1454

RESULT 8
A46269
sodium channel alpha chain HBA - human
C:Species: Homo sapiens (man)
C>Date: 20-Oct-1993 #sequence_revision 18-Nov-1994 #text_change 21-Nov-1997
C:Accession: A46269
R:Almed, C.M.; Ware, D.H.; Lee, S.C.; Patten, C.D.; Ferrer-Montiel, A.V.; Schindler, A.F.
Proc. Natl. Acad. Sci. U.S.A. 89, 8220-8224, 1992
A:Title: Primary structure, chromosomal localization, and functional expression of a vol
A:Reference number: A46269; MUID:92390418; PMID:1335650
A:Accession: A46269
A:Molecule type: mRNA
A:Residues: 1-2005 <AHM>
A:Cross-references: GB:M94055
A:Experimental source: brain
A:Note: sequence extracted from NCBI backbone (NCBI:P113082)
C:Genetics:
A:Map position: 2q23-q24.3
C:Superfamily: sodium channel protein
C:Keywords: duplication

Query Match 100.0%; Score 28; DB 2; Length 2005;
Best Local Similarity 100.0%; Pred. No. 3.7e-20;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 MYLYFVFIIFGSPFTLNLFGVIIDNF 28

Db 1449 MYLFFVFIIFGSEFTLNLFIVIIDNF 1476

RESULT 9

B25019

sodium channel protein II - rat

C/Species: Rattus norvegicus (Norway rat)

C/Date: 30-Jun-1988 #sequence (revision 30-Jun-1988 #text_change 09-Jul-2004

C/Accession: B25019; S24804

R/Node: M.; Ikeda, T.; Kayano, T.; Suzuki, H.; Takeshima, H.; Kurasaki, M.; Takahashi, H.

A/Title: Existence of distinct sodium channel messenger RNAs in rat brain.

A/Reference number: A93377; MUID:86146901; PMID:3754035

A/Accession: B25019

A/Molecule type: mRNA

A/Residues: 1-2005 <NOD>

A/Cross-references: UNIPROT:Q63509

R/Sarao, R.; Gupta, S.K.; Auld, V.J.; Dunn, R.J.

Submitted to the EMBL Data Library, August 1991.

A/Description: Developmentally regulated RNA splicing of rat brain sodium channel mRNAs.

A/Reference number: S24803

A/Accession: S24804

A/Status: preliminary

A/Molecule type: DNA

A/Residues: 183-188, 'D', 190-305 <SNR>

A/Cross-references: EMBL:X61149; NID:957074; PIDN:CAA43458.1; PID:957076

C/Suprafamily: sodium channel protein

C/Keywords: duplication; ion transport; sodium channel; transmembrane protein; voltage-g

Query Match

Best Local Similarity 100.0%; Score 28; DB 2; Length 2005;

Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MYLFFVFIIFGSEFTLNLFIVIIDNF 28

Db 1449 MYLFFVFIIFGSEFTLNLFIVIIDNF 1476

RESULT 10
A25019
sodium channel protein I - rat

N/Alternate names: sodium channel protein A

C/Species: Rattus norvegicus (Norway rat)

C/Date: 30-Jun-1988 #sequence (revision 30-Jun-1988 #text_change 09-Jul-2004

C/Accession: A25019; S40783; I64764

R/Node: M.; Ikeda, T.; Kayano, T.; Suzuki, H.; Takeshima, H.; Kurasaki, M.; Takahashi, H.

A/Title: Existence of distinct sodium channel messenger RNAs in rat brain.

A/Reference number: A93377; MUID:86146901; PMID:3754035

A/Accession: A25019

A/Molecule type: mRNA

A/Residues: 1-2009 <NOD>

A/Cross-references: UNIPROT:P04774; GB:X03638; NID:957216; PIDN:CAA27286.1; PID:957217

R/Sarao, R.; Gupta, S.K.; Auld, V.J.; Dunn, R.J.

Nucleic Acids Res. 19, 5673-5679, 1991.

A/Title: Developmentally regulated alternative RNA splicing of rat brain sodium channel

A/Reference number: S40782; MUID:92051314; PMID:1658739

A/Accession: S40783

A/Molecule type: DNA

A/Residues: 177-253 <SNR>

R/Node: M.; Numa, S.

J. Recept. Res. 7, 467-497, 1987

A/Title: Structure and function of sodium channel.

A/Reference number: 150536; MUID:87311395; PMID:2442385

A/Accession: 184764

A/Status: preliminary; translated from GB/EMBL/DBJ

A/Residues: 1-2009 <RES>

A/Cross-references: GB:M22253; NID:91041088; PIDN:AAA9965.1; PID:91041089

C/Suprafamily: sodium channel protein

C/Keywords: duplication; ion transport; sodium channel; transmembrane protein; voltage-g

Query Match
Best Local Similarity 100.0%; Score 28; DB 2; Length 2009;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MYLFFVFIIFGSEFTLNLFIVIIDNF 28

Db 1459 MYLFFVFIIFGSEFTLNLFIVIIDNF 1486

RESULT 11

I56555

sodium channel protein 6 - rat

C/Species: Rattus norvegicus (Norway rat)

C/Date: 26-Jul-1996 #sequence (revision 26-Jul-1996 #text_change 09-Jul-2004

C/Accession: I56555

R/Schaller, K.L.; Krzemien, D.M.; Yarowsky, P.J.; Krueger, B.K.; Caldwell, J.H.

J. Neurosci. 15, 3231-3242, 1995

A/Title: A novel, abundant sodium channel expressed in neurons and glia.

A/Reference number: I56555; MUID:95271284; PMID:7751906

A/Accession: I56555

A/Status: preliminary; translated from GB/EMBL/DBJ

A/Molecule type: mRNA

A/Residues: 1-1976 <RES>

A/Cross-references: UNIPROT:Q63541; GB:I39018; NID:9829033; PIDN:AA42059.1; PID:9829034

C/Suprafamily: sodium channel protein

C/Keywords: duplication

Query Match
Best Local Similarity 89.3%; Score 25; DB 2; Length 1976;

Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 YFVFIIFGSEFTLNLFIVIIDNF 28

Db 1439 YFVFIIFGSEFTLNLFIVIIDNF 1463

RESULT 12
A38195
sodium channel protein hH1, cardiac - human

N/Alternate names: tetrodotoxin-insensitive, voltage-dependent sodium channel, TTX-I Na

C/Species: Homo sapiens (man)

C/Date: 31-Dec-1993 #sequence (revision 31-Dec-1993 #text_change 09-Jul-2004

C/Accession: A38195

R/Gellens, M.E.; George Jr., A.L.; Chen, L.O.; Chahine, M.; Horn, R.; Barchi, R.L.; Kall

Proc. Natl. Acad. Sci. U.S.A. 89, 554-558, 1992

A/Title: Primary structure and functional expression of the human cardiac tetrodotoxin-1

A/Reference number: A38195; MUID:92115699; PMID:1309946

A/Accession: A38195

A/Status: nucleic acid sequence not shown

A/Molecule type: mRNA

A/Residues: 1-2016 <GEL>

A/Cross-references: UNIPROT:Q14524; GB:M77235; NID:9184038; PIDN:AAA58644.1; PID:9184039

R/Schaller, K.L.; Krzemien, D.M.; Yarowsky, P.J.; Krueger, B.K.; Caldwell, J.H.

J. Neurosci. 15, 3231-3242, 1995

A/Title: A novel, abundant sodium channel expressed in neurons and glia.

A/Reference number: I56555; MUID:95271284; PMID:7751906

A/Accession: I56555

A/Status: preliminary; translated from GB/EMBL/DBJ

A/Molecule type: mRNA

A/Residues: 1-1976 <RES>

A/Cross-references: UNIPROT:Q63541; GB:I39018; NID:9829033; PIDN:AA42059.1; PID:9829034

C/Suprafamily: sodium channel protein

C/Keywords: cardiac muscle; duplication; glycoprotein; heart; ion transport; sodium chan

Query Match

Best Local Similarity 89.3%; Score 25; DB 2; Length 2016;

Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 YFVFIIFGSEFTLNLFIVIIDNF 28

Db 1449 YFVFIIFGSEFTLNLFIVIIDNF 1473

RESULT 13
A56595
sodium channel alpha chain hscp - tobacco budworm (fragment)

C/Species: Heliothis virescens (tobacco budworm)

C/Date: 11-Aug-1995 #sequence (revision 11-Aug-1995 #text_change 09-Jul-2004

C/Accession: A56595
 R/Taylor, M.F.; Heckel, D.G.; Brown, T.M.; Kretzman, M.E.; Black, B.
 Insect Biochem. Mol. Biol. 23, 763-775, 1993
 A/Title: Linkage of pyrethroid insecticide resistance to a sodium channel locus in the C
 A/Reference number: A56595; MUID:93386183; PMID:8397035
 A/Accession: A56595
 A/Status: preliminary
 A/Molecule type: DNA
 A/Residues: 1-295 <TAY>
 A/Cross-references: UNIPROT:Q25178; GB:U24236
 A/Experimental source: pyrethroid resistant strain PEG-87
 A/Note: authors translated the codons GGC for residue 122 as Ala, GTA for residue 238 as
 C/Superfamily: sodium channel protein
 C/Keywords: duplication

Query Match 75.0%; Score 21; DB 2; Length 295;
 Best Local Similarity 100.0%; Pred. No. 7.2e-14;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 8 FIIFGSFPTNLFIGVIIDNF 28
 Db 94 FIIFGSFPTNLFIGVIIDNF 114

RESULT 14
 S72467
 sodium channel protein para-type alpha chain - German cockroach (strain CSMA) (fragment)
 C/Species: Blattella germanica (German cockroach)
 A/Variety: strain CSMA
 C/Date: 29-Jul-1997 #sequence_revision 29-Aug-1997 #text_change 09-Jul-2004
 C/Accession: S72467; S72487
 R/Miyazaki, M.; Ohyama, K.; Dunlap, D.Y.; Matsunura, F.
 submitted to the EMBL Data Library, September 1996
 A/Description: Cloning and sequencing of the para-type sodium channel gene from suscepti
 A/Reference number: S72467
 A/Accession: S72467
 A/Molecule type: mRNA
 A/Residues: 1-1689 <MIY>
 A/Cross-references: UNIPROT:Q93135; EMBL:U71083; NID:g1633647; PIDN:AB82037.1; PID:g163
 Mol. Gen. Genet. 252, 61-68, 1996
 A/Title: Cloning and sequencing of the para-type sodium channel gene from susceptible ar
 A/Reference number: S72487; MUID:96397510; PMID:8804404
 A/Accession: S72487
 A/Molecule type: mRNA
 A/Residues: 711-819 <MIW>
 A/Cross-references: EMBL:U71083
 C/Superfamily: sodium channel protein
 C/Keywords: duplication; sodium channel; transmembrane protein

Query Match 75.0%; Score 21; DB 2; Length 1689;
 Best Local Similarity 100.0%; Pred. No. 3.3e-13;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 8 FIIFGSFPTNLFIGVIIDNF 28
 Db 1304 FIIFGSFPTNLFIGVIIDNF 1324

RESULT 15
 T31340
 voltage-gated sodium channel homolog - Bde11oura candida
 C/Species: Bde11oura candida
 C/Date: 02-Sep-2000 #sequence_revision 02-Sep-2000 #text_change 09-Jul-2004
 C/Accession: T31340
 R/Jezioraki, M.C.; Greenberg, R.M.; Anderson, P.A.
 submitted to the EMBL Data Library, March 1997
 A/Description: A putative voltage-gated sodium channel from the turbellarian flatworm Bd
 A/Reference number: Z21006
 A/Accession: T31340
 A/Status: preliminary; translated from GB/EMBL/DBJ
 A/Molecule type: mRNA
 A/Residues: 1-1699 <JEZ>

A/Cross-references: UNIPROT:O02037; EMBL:U93074; NID:g1947093; PID:g1947094; PIDN:AAC630
 C/Genetics:
 A/Gene: Nat1
 C/Superfamily: sodium channel protein

Query Match 75.0%; Score 21; DB 2; Length 1699;
 Best Local Similarity 100.0%; Pred. No. 3.3e-13;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 8 FIIFGSFPTNLFIGVIIDNF 28
 Db 1231 FIIFGSFPTNLFIGVIIDNF 1251

RESULT 16
 T42388
 sodium channel alpha chain - rat
 C/Species: Rattus norvegicus (Norway rat)
 C/Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 09-Jul-2004
 C/Accession: T42388
 R/Did-Hajj, S.D.; Tyrrell, L.; Black, J.A.; Waxman, S.G.
 Proc. Natl. Acad. Sci. U.S.A. 95, 8963-8968, 1998
 A/Title: NaN, a novel voltage-gated Na channel, is expressed preferentially in periphera
 A/Reference number: Z22149; MUID:98338024; PMID:9671787
 A/Accession: T42388
 A/Status: preliminary; translated from GB/EMBL/DBJ
 A/Molecule type: mRNA
 A/Residues: 1-1765 <DIB>
 A/Cross-references: UNIPROT:O88457; EMBL:AF059030; NID:g3372614; PID:g3372615; PIDN:AAC4
 A/Experimental source: strain Sprague-Dawley; dorsal root ganglia
 A/Note: preferentially expressed in sensory neurons within dorsal root ganglia and trigem
 C/Superfamily: sodium channel protein

Query Match 75.0%; Score 21; DB 2; Length 1765;
 Best Local Similarity 100.0%; Pred. No. 3.4e-13;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 8 FIIFGSFPTNLFIGVIIDNF 28
 Db 1271 FIIFGSFPTNLFIGVIIDNF 1291

RESULT 17
 T43167
 sodium channel protein - California market squid
 C/Species: Loligo opalescens (California market squid)
 C/Date: 11-Jan-2000 #sequence_revision 11-Jan-2000 #text_change 09-Jul-2004
 C/Accession: T43167
 R/Rosenthal, J.J.; Gilly, W.F.
 Proc. Natl. Acad. Sci. U.S.A. 90, 10026-10030, 1993
 A/Title: Amino acid sequence of a putative sodium channel expressed in the giant axon of
 A/Reference number: Z22324; MUID:94052096; PMID:8234251
 A/Accession: T43167
 A/Status: preliminary; translated from GB/EMBL/DBJ
 A/Residues: 1-1784 <ROS>
 A/Molecule type: mRNA
 A/Cross-references: UNIPROT:Q25377; EMBL:L19979; NID:g349118; PID:g349119; PIDN:AAA16202
 A/Experimental source: stellate ganglia
 C/Superfamily: sodium channel protein
 C/Keywords: ion transport; membrane protein; sodium channel; voltage-gated ion channel

Query Match 75.0%; Score 21; DB 2; Length 1784;
 Best Local Similarity 100.0%; Pred. No. 3.4e-13;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 8 FIIFGSFPTNLFIGVIIDNF 28
 Db 1268 FIIFGSFPTNLFIGVIIDNF 1288

RESULT 18
 A33299
 sodium channel protein - fruit fly (Drosophila melanogaster) (fragment)

C:Species: Drosophila melanogaster
C:Date: 20-Dec-1989 #sequence_revision 20-Dec-1989 #text_change 21-Nov-1997
C:Accession: A33299
R:Loughney, K.; Kreber, R.; Ganetzky, B.
Cell 58, 1143-1154, 1989
A:Title: Molecular analysis of the para locus, a sodium channel gene in Drosophila.
A:Reference number: A33299; PMID:89376565; PMID:2550145
A:Accession: A33299
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-1820 <IDU>
A:Cross-references: GB:M2078; GB:M24285
C:Genetics:
A:Gene: FlyBase:para
A:Cross-references: FlyBase:FBgn0003036
C:Superfamily: sodium channel protein
C:Keywords: duplication; phosphoprotein; transmembrane protein

Query Match
Best Local Similarity 75.0%; Score 21; DB 2; Length 1820;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 FIIFGSFPTLNLFIGVITDNF 28
DB 1499 FIIFGSFPTLNLFIGVITDNF 1519

RESULT 19
S54771
sodium channel alpha subunit - human
C:Species: Homo sapiens (man)
C:Date: 27-Oct-1995 #sequence_revision 03-Nov-1995 #text_change 09-Jul-2004
C:Accession: S54771
R:Kupfer, N.; Lacinova, L.; Flockerzi, V.; Hofmann, F.
EMBO J. 14, 1084-1090, 1995
A:Title: Structure and functional expression of a new member of the tetrodotoxin-sensitive
A:Reference number: S54771; PMID:95237189; PMID:7720699
A:Accession: S54771
A:Status: preliminary; nucleic acid sequence not shown
A:Molecule type: mRNA
A:Residues: 1-1977 <KLU>
A:Cross-references: UNIPROT:Q15858; EMBL:X82835; NID:9758109; PIDN:GMS8042.1; PID:97581
C:Superfamily: sodium channel protein
C:Keywords: duplication

Query Match
Best Local Similarity 75.0%; Score 21; DB 2; Length 1977;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 FIIFGSFPTLNLFIGVITDNF 28
DB 1429 FIIFGSFPTLNLFIGVITDNF 1449

RESULT 20
A33996
sodium channel protein I, cardiac - rat
N:Alternate names: sodium channel protein (SKM2) alpha chain
C:Species: Rattus norvegicus (Norway rat)
C:Date: 30-Mar-1990 #sequence_revision 30-Mar-1990 #text_change 09-Jul-2004
C:Accession: A33996; J00412
R:Rogart, R.B.; Cribbs, L.L.; Muglia, L.K.; Kephart, D.D.; Kaiser, M.W.
Proc. Natl. Acad. Sci. U.S.A. 86, 8170-8174, 1989
A:Title: Molecular cloning of a putative tetrodotoxin-resistant rat heart Na(+) channel
A:Reference number: A33996; PMID:90046760; PMID:25543032
A:Accession: A33996
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-2019 <ROG>
A:Cross-references: UNIPROT:P15389; GB:M27902; NID:9206857; PIDN:AAA42114.1; PID:9206858
R:Kallen, R.G.; Sheng, Z.H.; Yang, J.; Chen, L.; Rogart, R.B.; Barchi, R.L.
Neuron 4, 233-242, 1990
A:Title: Primary structure and expression of a sodium channel characteristic of denervat

A:Reference number: J00412; PMID:90166613; PMID:2155010
A:Accession: J00412
A:Molecule type: mRNA
A:Residues: 1-479, 481-1712, 'T', 1714-1963, 'R', 1965-2019 <RAL>
A:Experimental source: muscle
C:Superfamily: sodium channel protein
C:Keywords: cardiac muscle; duplication; heart; sodium channel; transmembrane protein

Query Match
Best Local Similarity 75.0%; Score 21; DB 2; Length 2019;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 FIIFGSFPTLNLFIGVITDNF 28
DB 1455 FIIFGSFPTLNLFIGVITDNF 1475

RESULT 21
S72458
sodium channel protein para-type alpha chain - house fly (strain Cooper)
C:Species: Musca domestica (house fly)
A:Variety: strain Cooper
C:Date: 24-Oct-1998 #sequence_revision 24-Oct-1998 #text_change 09-Jul-2004
C:Accession: S72458
R:Williamson, M.S.; Martinez-Torres, D.; Hick, C.A.; Devonshire, A.L.
Mol. Gen. Genet. 252, 51-60, 1996
A:Title: Identification of mutations in the housefly para-type sodium channel gene associ
A:Reference number: S72458; PMID:96397509; PMID:8804403
A:Accession: S72458
A:Molecule type: mRNA
A:Residues: 1-2108 <WIL>
A:Cross-references: UNIPROT:Q94615; EMBL:X96668
A:Experimental source: strain Cooper
C:Genetics:
A:Map position: 3
C:Superfamily: sodium channel protein
C:Keywords: alternative splicing; glycoprotein; phosphoprotein; sodium channel; transmem
F:302, 314, 332, 967, 1451, 1470/binding site: carbohydrate (asn) (covalent) #status predicted
F:541, 1208, 1582/binding site: phosphate (ser) (covalent) #status predicted

Query Match
Best Local Similarity 75.0%; Score 21; DB 2; Length 2108;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 FIIFGSFPTLNLFIGVITDNF 28
DB 1531 FIIFGSFPTLNLFIGVITDNF 1551

RESULT 22
T43161
sodium channel protein Tuna1 - sea squirt (Halocynthia roretzi)
C:Species: Halocynthia roretzi
C:Date: 11-Jan-2000 #sequence_revision 11-Jan-2000 #text_change 09-Jul-2004
C:Accession: T43161
R:Okamura, Y.; Ono, F.; Okagaki, R.; Chong, J.; Mandel, G.
Neuron 13, 937-948, 1994
A:Title: Neutral expression of a sodium channel gene requires cell-specific interactions.
A:Reference number: Z22220; PMID:95033215; PMID:7946338
A:Accession: T43161
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-2049 <OKA>
A:Cross-references: UNIPROT:Q25150; EMBL:DJ7311; PIDN:BAA04133.1
C:Superfamily: sodium channel protein
C:Keywords: sodium channel; transmembrane protein

Query Match
Best Local Similarity 64.3%; Score 18; DB 2; Length 2049;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 11 FGSGFTLNLFIGVITDNF 28
|||||

DB 1522 GSFPTLNLFIGVIIDNF 1539

RESULT 23

A60165 sodium channel protein - fruit fly (*Drosophila melanogaster*) (fragments)

C/Species: *Drosophila melanogaster*

C/Date: 22-Jan-1993 #sequence_revision 22-Jan-1993 #text_change 09-Jul-2004

C/Accession: S04029; A60165

R/Salkoff, L.; Butler, A.; Scavarda, N.; Wei, A.

Nucleic Acids Res. 15, 8569-8572, 1987

A/Title: Nucleotide sequence of the putative sodium channel gene from *Drosophila*: the fo

A/Reference number: S04029; MUID:88040482; PMID:2444928

A/Accession: S04029

A/Molecule type: DNA

A/Residues: 1362;363-626;627-1321 <SNL>

A/Cross-references: UNIPROT:Q27930; EMBL:X14394

R/Salkoff, L.; Butler, A.; Wei, A.; Scavarda, N.; Giffen, K.; Ifune, C.; Goodman, R.; Ma

Science 237, 744-749, 1987

A/Title: Genomic organization and deduced amino acid sequence of a putative sodium chan

A/Reference number: A60165; MUID:87292090; PMID:2441469

A/Accession: A60165

A/Status: nucleic acid sequence not shown

A/Molecule type: DNA

A/Residues: 40-355;363-560, 'P', 562-626;632-1263 <SA2>

A/Cross-references: EMBL:X14394

A/Note: part of this sequence was confirmed by mRNA sequencing

A/Note: the authors' translation is shown at position 561

C/Genetics:

A/Genes: *FlyBase:NACP60E*

A/Cross-references: *FlyBase:Fggn0002920*

A/Introns: 237/2; 310/3; 362/3; 414/3; 471/3; 531/3; 581/1; 626/3; 751/2; 801/1; 908/1;

Query Match 60.7%; Score 17; DB 2; Length 1321;

Best Local Similarity 100.0%; Pred. No. 2.7e-09;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 12 GSFPTLNLFIGVIIDNF 28

DB 886 GSFPTLNLFIGVIIDNF 902

RESULT 24

T31092

probable voltage-gated sodium channel - *Aiptasia pallida*

C/Species: *Aiptasia pallida*

C/Date: 02-Sep-2000 #sequence_revision 02-Sep-2000 #text_change 09-Jul-2004

C/Accession: T31092

R/White, G.B.; Pfahnl, A.; Haddock, S.; Lamers, S.; Greenberg, R.M.; Anderson, P.A.V.

submitted to the EMBL Data Library, January 1998

A/Description: Structure of a putative sodium channel from the sea anemone *Aiptasia pall*

A/Reference number: Z20975

A/Accession: T31092

A/Status: preliminary; translated from GB/EMBL/DBJ

A/Molecule type: mRNA

A/Residues: 1-1810 <MHI>

A/Cross-references: UNIPROT:O44930; EMBL:AF041851; NID:g2791840; PID:g2791841; PIDN:AA85

C/Genetics:

A/Genes: *Nal*

C/Superfamily: sodium channel protein

Query Match 60.7%; Score 17; DB 2; Length 1810;

Best Local Similarity 100.0%; Pred. No. 3.5e-09;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 12 GSFPTLNLFIGVIIDNF 28

DB 1294 GSFPTLNLFIGVIIDNF 1310

RESULT 25

CHBE

sodium channel protein - electric eel

C/Species: *Electrophorus electricus* (electric eel)

C/Date: 28-May-1986 #sequence_revision 28-May-1986 #text_change 09-Jul-2004

C/Accession: A03178; I50536

R/Noda, M.; Shimizu, S.; Tanabe, T.; Takai, T.; Kayano, T.; Ikeda, T.; Takahashi, H.; Na

da, H.; Miyata, T.; Numa, S.

Nature 312, 121-127, 1984

A/Title: Primary structure of *Electrophorus electricus* sodium channel deduced from cDNA

A/Reference number: A03178; MUID:85061498; PMID:6209577

A/Accession: A03178

A/Molecule type: mRNA

A/Residues: 1-1820 <MOD>

A/Cross-references: UNIPROT:P02719; GB:X01119; NID:962776; PIDN:CAA25587.1; PID:962777

R/Noda, M.; Numa, S.

J. Recept. Res. 7, 467-497, 1987

A/Title: Structure and function of sodium channel.

A/Reference number: I50536; MUID:87311395; PMID:2442385

A/Accession: I50536

A/Status: preliminary; translated from GB/EMBL/DBJ

A/Molecule type: mRNA

A/Residues: 1-1820 <MOD>

A/Cross-references: GB:M22252; NID:91041048; PIDN:AA79960.1; PID:91041049

C/Comment: This membrane glycoprotein mediates the voltage-dependent sodium-ion permeabil

as the membrane, the protein forms a sodium-selective channel through which sodium ions m

C/Comment: This sequence contains four highly homologous internal repeats (excluding res

Each repeat has a similar overall structure containing six subregions located in identic

has a net positive charge (S4), and one is neutral (S2).

C/Comment: The four repeating units are thought to be oriented pseudosymmetrically across

e. The presence of four homologous structures within this molecule is consistent with the

C/Comment: Available data suggest that activation and inactivation gates are located near

955 might, in conjunction with the positively charged residues of S4, act as a voltage se

C/Superfamily: sodium channel protein

A/KeyWords: duplication; glycoprotein; ion transport; membrane protein; sodium channel; \

F:111-419,555-585,989-1019,1311-1341/Region: S1 of repeats I through IV

F:150-171,597-620,1033-1057,1353-1376/Region: S2 of repeats I through IV

F:177-197,626-643,1062-1079,1381-1398/Region: S3 of repeats I through IV

F:204-224,651-671,1092-1112,1417-1437/Region: S4 of repeats I through IV

F:244-264,691-711,1132-1152,1454-1474/Region: S5 of repeats I through IV

F:379-402,767-790,1236-1264,1544-1567/Region: S6 of repeats I through IV

F:205,278,288,317,591,690,797,1160,1174,1806/Binding site: carbohydrate (asn) (covalent)

Query Match 53.6%; Score 15; DB 1; Length 1820;

Best Local Similarity 100.0%; Pred. No. 3.6e-07;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 14 FPTLNLFIGVIIDNF 28

DB 1253 FPTLNLFIGVIIDNF 1267

Search completed: January 27, 2005, 17:52:46
Job time : 19 secs

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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 27, 2005, 17:35:20 ; Search time 92.5 Seconds
(without alignment)
174.167 Million cell updates/sec

Title: US-10-608-584-19
Perfect score: 28
Sequence: 1 MYLVFVIFIFGSPFTLNLFIGVIIDNF 28

Scoring table: OLIGO
Gapop 60.0 , Gapext 60.0

Searched: 1825181 seqs, 575374646 residues

Word size : 0

Total number of hits satisfying chosen parameters: 1825181

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Listing first 100 summaries

Database : Uniprot_02.*
1: uniprot_sprot.*
2: uniprot_trembl.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	28	100.0	431	2	Q62206
2	28	100.0	510	2	Q62242
3	28	100.0	742	2	Q90229
4	28	100.0	751	2	Q90228
5	28	100.0	1136	2	Q804F4
6	28	100.0	1836	1	CIN4_HUMAN
7	28	100.0	1840	1	CIN4_RAT
8	28	100.0	1840	2	Q70611
9	28	100.0	1841	2	Q9ER60
10	28	100.0	1951	1	CIN3_RAT
11	28	100.0	1951	2	Q9C007
12	28	100.0	1981	2	Q81U06
13	28	100.0	2000	1	CIN3_HUMAN
14	28	100.0	2005	1	CIN2_HUMAN
15	28	100.0	2005	1	CIN2_RAT
16	28	100.0	2007	2	Q9YGN7
17	28	100.0	2009	1	CIN1_HUMAN
18	28	100.0	2009	1	CIN1_RAT
19	28	89.3	718	2	Q90230
20	25	89.3	743	2	Q90226
21	25	89.3	1880	2	Q91BR1
22	25	89.3	1949	2	Q9DF53
23	25	89.3	1962	2	Q75RX9
24	25	89.3	1962	2	BAD12085
25	25	89.3	1976	2	Q63541
26	25	89.3	1976	1	CIN8_MOUSE
27	25	89.3	1978	2	Q884Z0
28	25	89.3	1980	1	CIN8_HUMAN
29	25	89.3	1984	2	Q28644
30	25	89.3	1984	2	Q08562
31	25	89.3	1988	2	Q88421

32	25	89.3	2013	2	Q865W3	Q865W3 canis famil
33	25	89.3	2015	2	Q86UR3	Q86UR3 homo sapien
34	25	89.3	2015	2	Q81ZC9	Q81ZC9 homo sapien
35	25	89.3	2015	2	Q96J69	Q96J69 homo sapien
36	25	89.3	2016	1	CIN5_HUMAN	Q14524 homo sapien
37	25	89.3	2016	2	Q75RY0	Q75RY0 homo sapien
38	25	89.3	2016	2	BAD12084	BAD12084 homo sapi
39	25	89.3	2022	2	Q8WMP8	Q8WMP8 bos taurus
40	21	75.0	169	2	Q7PJH1	Q7PJH1 anopheles g
41	21	75.0	210	2	Q9JRC5	Q9JRC5 mus musculu
42	21	75.0	296	2	Q25178	Q25178 heliothis v
43	21	75.0	576	2	Q6DLU1	Q6DLU1 aedes aegy
44	21	75.0	603	2	Q6DLT6	Q6DLT6 aedes albop
45	21	75.0	626	2	Q6DLT5	Q6DLT5 aedes albop
46	21	75.0	744	2	Q90Z27	Q90Z27 sternopygus
47	21	75.0	1130	2	Q9XZC1	Q9XZC1 boophilus m
48	21	75.0	1347	2	Q7PMT4	Q7PMT4 anopheles g
49	21	75.0	1361	2	Q7PJH0	Q7PJH0 anopheles g
50	21	75.0	1444	2	Q9JHMO	Q9JHMO homo sapien
51	21	75.0	1689	2	Q93135	Q93135 blatella g
52	21	75.0	1695	2	Q94584	Q94584 heliothis v
53	21	75.0	1699	2	Q02037	Q02037 bdelloura c
54	21	75.0	1717	2	Q90519	Q90519 figu rubrip
55	21	75.0	1765	2	Q88457	Q88457 rattus norv
56	21	75.0	1765	2	Q9JMD4	Q9JMD4 mus musculu
57	21	75.0	1765	2	Q9R053	Q9R053 mus musculu
58	21	75.0	1784	2	Q25377	Q25377 loligo opal
59	21	75.0	1791	2	Q8NDX3	Q8NDX3 homo sapien
60	21	75.0	1791	2	Q9JHE0	Q9JHE0 homo sapien
61	21	75.0	1791	2	Q9JH33	Q9JH33 homo sapien
62	21	75.0	1966	2	Q92566	Q92566 rattus norv
63	21	75.0	1977	2	Q15858	Q15858 homo sapien
64	21	75.0	2019	1	CIN5_RAT	P15389 rattus norv
65	21	75.0	2019	2	Q9JTV9	Q9JTV9 mus musculu
66	21	75.0	2031	2	Q01306	Q01306 blatella g
67	21	75.0	2031	2	Q01307	Q01307 blatella g
68	21	75.0	2051	2	Q86DI7	Q86DI7 pediculus h
69	21	75.0	2051	2	Q86DI8	Q86DI8 pediculus h
70	21	75.0	2051	2	Q86DI9	Q86DI9 pediculus h
71	21	75.0	2058	2	Q6DLT4	Q6DLT4 aedes albop
72	21	75.0	2064	2	Q6DLT3	Q6DLT3 aedes aegy
73	21	75.0	2086	2	Q86M38	Q86M38 pediculus h
74	21	75.0	2104	2	Q25440	Q25440 musca domes
75	21	75.0	2105	2	Q25439	Q25439 musca domes
76	21	75.0	2108	2	Q94615	Q94615 musca domes
77	21	75.0	2131	1	CIN4_DROME	P35500 drosophila
78	18	64.3	2049	2	Q25150	Q25150 halocynthia
79	17	60.7	281	2	Q7JN87	Q7JN87 drosophila
80	17	60.7	297	2	Q6VVE0	Q6VVE0 hirtudo medi
81	17	60.7	297	2	AAQ81289	AAQ81289 hirtudo me
82	17	60.7	1538	2	Q7PF76	Q7PF76 anopheles g
83	17	60.7	1618	2	Q8MWC7	Q8MWC7 drosophila
84	17	60.7	1810	2	Q44930	Q44930 alptasia pa
85	17	60.7	2223	2	Q7Q1V0	Q7Q1V0 anopheles g
86	17	60.7	2304	2	Q9BMO4	Q9BMO4 blatella g
87	17	60.7	2327	2	Q9W0Y8	Q9W0Y8 drosophila
88	16	57.1	1834	2	Q28371	Q28371 equus cabal
89	15	53.6	299	2	Q6VVE1	Q6VVE1 hirtudo medi
90	15	53.6	299	2	AAQ81288	AAQ81288 hirtudo me
91	15	53.6	301	2	Q6VVE2	Q6VVE2 hirtudo medi
92	15	53.6	301	2	AAQ81287	AAQ81287 hirtudo me
93	15	53.6	1089	2	Q81S97	Q81S97 varroa dest
94	15	53.6	1820	1	CIN4_ELEBL	P02719 electrophor
95	15	53.6	2215	2	Q86D77	Q86D77 varroa dest
96	13	46.4	1993	2	Q90670	Q90670 apllysia cal
97	12	42.9	717	2	Q90Z25	Q90Z25 sternopygus
98	12	42.9	1695	2	Q62604	Q62604 polyorchis
99	12	42.9	1740	2	Q17314	Q17314 cyanea capi
100	12	42.9	1956	2	Q9Y5Y9	Q9Y5Y9 homo sapien

ALIGNMENTS

RESULT 1

```

Q62206 ID Q62206 PRELIMINARY; PRT; 431 AA.
AC Q62206;
DR 01-NOV-1996 (TRENBLREL. 01, Created)
DT 01-NOV-1996 (TRENBLREL. 01, Last sequence update)
DE 01-MAR-2004 (TRENBLREL. 26, Last annotation update)
DR Sodium channel 3.
GN Name=Scnla;
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Brain;
RA Jover E., Shah V.;
RU Submitted (MAY-1995) to the EMBL/GenBank/DBJ databases.
DR EMBL; LA2339; AAA67107.1; -.
DR HSSP; P04775; 1BYV.
DR MGD; MGI:98246; Scnla.
DR GO; GO:0005624; C:membrane fraction; IDA.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR002111; Cat_channel_TrypL.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR005820; M+channel_nlg.
DR Pfam; PF00520; Ion_trans_1.
DR Pfam; PF06512; Na_trans_assoc; 1.
KW Ion transport; Ionic channel; Transmembrane; Transport.
SQ SEQUENCE 431 AA; 4946 MW; 9CD841FF73D1D9B7 CRC64;

Query Match 100.0%; Score 28; DB 2; Length 431;
Best Local Similarity 100.0%; Pred. No. 1.6e-19;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MYLVEVFIIFGSPFTLNLFIVGIIDNF 28
Db 357 MYLVEVFIIFGSPFTLNLFIVGIIDNF 384

RESULT 2
Q62242 ID Q62242 PRELIMINARY; PRT; 510 AA.
AC Q62242;
DR 01-NOV-1996 (TRENBLREL. 01, Created)
DT 01-NOV-1996 (TRENBLREL. 01, Last sequence update)
DE 01-MAR-2004 (TRENBLREL. 26, Last annotation update)
DR Sodium channel (Fragment).
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Brain;
RA Jover E., Shah V.;
RU Submitted (JUN-1995) to the EMBL/GenBank/DBJ databases.
CC -1- SIMILARITY: Belongs to the sodium channel family.
DR EMBL; LA2341; AAA67695.1; -.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0001518; C:membrane; IEA.
DR GO; GO:0005261; F:cation-gated sodium channel complex; IEA.
DR GO; GO:0005248; F:voltage-gated sodium channel activity; IEA.
DR GO; GO:0006812; P:cation transport; IEA.
DR GO; GO:0006814; P:sodium ion transport; IEA.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR002111; Cat_channel_TrypL.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IQ_region.
DE InterPro; IPR005820; M+channel_nlg.

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DR InterPro; IPR001696; Na_channel.
DR Pfam; PF00520; Ion_trans; 1.
DR Pfam; PF00612; IQ; 1.
DR PRINTS; PR00170; NACHANNEL.
DR SMART; SM00015; IQ; 1.
DR PROSITE; PS50096; IQ; 1.
KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
Transport; Voltage-gated channel.
FT NON_TER 510
SQ SEQUENCE 510 AA; 58397 MW; 02DCC7DABD3796E8 CRC64;

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Query Match 100.0%; Score 28; DB 2; Length 510;
Best Local Similarity 100.0%; Pred. No. 1.8e-19;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MYLVEVFIIFGSPFTLNLFIVGIIDNF 28
Db 1 MYLVEVFIIFGSPFTLNLFIVGIIDNF 28

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RESULT 3

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Q90229 ID Q90229 PRELIMINARY; PRT; 742 AA.
AC Q90229;
DR 01-DEC-2001 (TRENBLREL. 19, Created)
DT 01-DEC-2001 (TRENBLREL. 19, Last sequence update)
DE 01-MAR-2004 (TRENBLREL. 26, Last annotation update)
DR Sodium channel 2 (Fragment).
OS Sternopygus macrurus.
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Osteichthyes; Gymnotiformes;
OC Sternopygidae; Sternopygus.
NCBI_TaxID=77841;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=21310016; PubMed=11416226;
RA Lopreato G.F., Lu Y., Southwell A., Atkinson N.S., Hillis D.M.,
RA Wilcox T.P., Zakon H.H.;
RT Evolution and divergence of sodium channel genes in vertebrates.
RL Proc. Natl. Acad. Sci. U.S.A. 98:7568-7592(2001).
DR EMBL; AF378140; AAK5438.1; -.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0005261; F:cation channel activity; IEA.
DR GO; GO:0006812; P:cation transport; IEA.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR002111; Cat_channel_TrypL.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR005820; M+channel_nlg.
DR Pfam; PF00520; Ion_trans_2.
DR Pfam; PF06512; Na_trans_assoc; 1.
KW Ion transport; Ionic channel; Transmembrane; Transport.
FT NON_TER 742
SQ SEQUENCE 742 AA; 84353 MW; 7F4B3003BD3F2AC4 CRC64;

Query Match 100.0%; Score 28; DB 2; Length 742;
Best Local Similarity 100.0%; Pred. No. 2.5e-19;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MYLVEVFIIFGSPFTLNLFIVGIIDNF 28
Db 641 MYLVEVFIIFGSPFTLNLFIVGIIDNF 668

RESULT 4
Q90228 ID Q90228 PRELIMINARY; PRT; 751 AA.
AC Q90228;
DR 01-DEC-2001 (TRENBLREL. 19, Created)
DT 01-DEC-2001 (TRENBLREL. 19, Last sequence update)
DE 01-MAR-2004 (TRENBLREL. 26, Last annotation update)
DR Sodium channel 3 (Fragment).

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OS Sternopygus macrurus.
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Osteiophysi; Gymnotiformes;
OC Sternopygidae; Sternopygus.
OX NCBI_TaxID=77841;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=21310016; PubMed=11416226;
RA Lopreato G.F., Lu Y., Southwell A., Atkinson N.S., Hillis D.M.,
RA Wilcox T.P., Zakon H.H.;
RT "Evolution and divergence of sodium channel genes in vertebrates.";
RL Proc. Natl. Acad. Sci. U.S.A. 98:7588-7592(2001).
DR EMBL; AF378141; AKS5439.1; -.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0005261; F:cation channel activity; IEA.
DR GO; GO:0006812; F:cation transport; IEA.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR002111; Cat_channel_TrypL.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR010526; Na_trans_assoc.
DR Pfam; PF00520; Ion_trans_2.
DR Pfam; PF06512; Na_trans_assoc_1.
KM Ion transport; Ionic channel; Transmembrane; Transport.
FT NON_TER 1
FT NON_TER 1
SQ SEQUENCE 751 AA; 84598 MW; CBF6162B90A76FC CRC64;

Query Match 100.0%; Score 28; DB 2; Length 751;
Best Local Similarity 100.0%; Pred. No. 2.5e-19;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 MYLYFVFIIFGSPFTLNLFIGVINDNF 28
Db 674 MYLYFVFIIFGSPFTLNLFIGVINDNF 701

RESULT 5
Q804F4 PRELIMINARY; PRT; 1136 AA.
ID Q804F4;
AC Q804F4;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE Sodium channel 7 (Fragment).
GN Name=Na7;
OS Sternopygus macrurus.
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Osteiophysi; Gymnotiformes;
OC Sternopygidae; Sternopygus.
OX NCBI_TaxID=77841;
RN [1]
RP SEQUENCE FROM N.A.
RA Lu Y., Lopreato G.F., Zakon H.H.;
RL Submitted (NOV-2002) to the EMBL/GenBank/DBJ databases.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
CC -1- SIMILARITY: Belongs to the calcium channel alpha-1 subunit
family.
DR EMBL; AY183895; AAO23570.1; -.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0005891; C:voltage-gated calcium channel complex; IEA.
DR GO; GO:0005261; F:cation channel activity; IEA.
DR GO; GO:0005245; F:voltage-gated calcium channel activity; IEA.
DR GO; GO:0006812; F:calcium ion transport; IEA.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR002111; Cat_channel_TrypL.
DR InterPro; IPR002077; Ca_channel_Alpha.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR005820; M_channel_nlg.
DR InterPro; IPR010526; Na_trans_assoc.
DR Pfam; PF00520; Ion_trans_3.
DR Pfam; PF06512; Na_trans_assoc_1.

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DR PRINTS; PR00167; CCHANNEL.
KM Calcium channel; Calcium-binding; Ion transport; Ionic channel;
KM Transmembrane; Transport; Voltage-gated channel.
FT NON_TER 1
FT NON_TER 1
SQ SEQUENCE 1136 AA; 129141 MW; ECD52D025B50664 CRC64;

Query Match 100.0%; Score 28; DB 2; Length 1136;
Best Local Similarity 100.0%; Pred. No. 3.5e-19;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 MYLYFVFIIFGSPFTLNLFIGVINDNF 28
Db 1042 MYLYFVFIIFGSPFTLNLFIGVINDNF 1069

RESULT 6
CIN4_HUMAN STANDARD; PRT; 1836 AA.
ID CIN4_HUMAN;
AC P35499; Q15478; Q16447; Q726B1;
DT 01-JUN-1994 (Rel. 29, Created)
DT 29-MAR-2004 (Rel. 43, Last sequence update)
DT 05-JUL-2004 (Rel. 44, Last annotation update)
DE Sodium channel protein type IV alpha subunit (Voltage-gated sodium
channel alpha subunit Nav1.4) (Sodium channel protein, skeletal muscle
DE alpha-subunit).
GN Name=SCN4A;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A., AND VARIANT ASP-559.
RC TISSUE=Skeletal muscle;
RX MEDLINE=92246457; PubMed=1315496;
RA George A.L. Jr., Komisarof J., Kallen R.G., Barchi R.L.;
RT "Primary structure of the adult human skeletal muscle voltage-
dependent sodium channel.";
RL Ann. Neurol. 31:131-137(1992).
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE=92134303; PubMed=1310396;
RA Wang J., Rojas C.V., Zhou J., Schwartz L.S., Nicholas H.,
RA Hoffmann E.P.;
RT "Sequence and genomic structure of the human adult skeletal muscle
sodium channel alpha subunit gene on 17q.";
RL Biochem. Biophys. Res. Commun. 182:794-801(1992).
RN [3]
RP SEQUENCE FROM N.A., VARIANT MYASTHENIC SYNDROME GLU-1442, AND VARIANTS
LEU-246; ASP-559 AND ASN-1376.
RX MEDLINE=22684480; PubMed=12762226; DOI=10.1073/pnas.1230273100;
RA Teujino A., Maertens C., Ohno K., Shen X.-M., Fukuda T., Harper C.M.,
RA Cannon S.C., Engel A.G.;
RT "Myasthenic syndrome caused by mutation of the SCN4A sodium channel.";
RL Proc. Natl. Acad. Sci. U.S.A. 100:7377-7382(2003).
RN [4]
RP SEQUENCE FROM N.A., AND VARIANT ASN-1376.
RX MEDLINE=9338444; PubMed=1339144;
RA McLatchey A.I., Ian C.S., Wang J., Hoffman E.P., Rojas C.V.,
RA Gusella J.F.;
RT "The genomic structure of the human skeletal muscle sodium channel
gene.";
RL Hum. Mol. Genet. 1:521-521(1992).
RN [5]
RP SEQUENCE OF 1305-1339 FROM N.A., AND VARIANTS PMC VAL-1306 AND
MET-1313.
RX MEDLINE=92154689; PubMed=1310898;
RA McLatchey A.I., van den Bergh P., Pericak-Vance M.A., Raskind W.,
RA Verellen C., McKenna-Yasek D., Rao K., Haines J.L., Bird T.,
RA Brown R.H. Jr., Gusella J.F.;
RT "Temperature-sensitive mutations in the III-IV cytoplasmic loop region
of the skeletal muscle sodium channel gene in paramyotonia
congenita.";

```

RL Cell 68:769-774(1992).
 RN [6]
 RP VARIANT HYP MET-704.
 RX MEDLINE=92069747; PubMed=1659948;
 RA Pracek L.J., George A.L. Jr., Griggs R.C., Tawil R., Kallen R.G.,
 RT "Identification of a mutation in the gene causing hyperkalemic
 RL periodic paralysis.";
 RN Cell 67:1021-1027(1991).
 RN [7]
 RP VARIANT HYP VAL-1592.
 RX MEDLINE=92065978; PubMed=1659668;
 RA Rojas C.V., Wang J., Schwartz U.S., Hoffman E.P., Powell B.R.,
 RT "A Met-1021 mutation in the skeletal muscle Na⁺ channel alpha-
 RL subunit in hyperkalemic periodic paralysis.";
 RN Nature 354:387-389(1991).
 RN [8]
 RP VARIANT PWC PHE-804 AND THR-1156.
 RX MEDLINE=93265141; PubMed=1338909;
 RA McGlatchey A.I., McKenna-Yasek D., Cros D., Worthen H.G., Kuncl R.W.,
 RT "Novel mutations in families with unusual and variable disorders of
 RL the skeletal muscle sodium channel.";
 RN Nat. Genet. 2:148-152(1992).
 RN [9]
 RP VARIANT PWC CYS-1448 AND HIS-1448.
 RX MEDLINE=92265302; PubMed=1316765;
 RA Pracek L.J., George A.L. Jr., Barohn R.L., Griggs R.C., Riggs J.E.,
 RT "Mutations in an S4 segment of the adult skeletal muscle sodium
 RL channel cause paramyotonia congenita.";
 RN Neuron 8:891-897(1992).
 RN [10]
 RP VARIANT PWC ARG-1433.
 RX MEDLINE=93270429; PubMed=8388676;
 RA Pracek L.J., Gow L., Kwleciniski H., McManis P., Mendell J.R.,
 RT "Sodium channel mutations in paramyotonia congenita and hyperkalemic
 RL periodic paralysis.";
 RN Ann. Neurol. 33:300-307(1993).
 RN [11]
 RP VARIANT PWC ALA-1306; GLU-1306 AND VAL-1306.
 RX MEDLINE=94141728; PubMed=8308722;
 RA Lerche H., Heine R., Pika U., George A.L. Jr., Mitrovic N.,
 RT "Human sodium channel myotonia: slowed channel inactivation due to
 RL substitutions for a glycine within the III-IV linker.";
 RN J. Physiol. (Lond.) 470:13-22(1993).
 RN [12]
 RP VARIANT PWC MET-1589.
 RX MEDLINE=94061027; PubMed=8242056;
 RA Heine R., Pika U., Lehmann-Horn F.,
 RT "A novel SCN4A mutation causing myotonia aggravated by cold and
 RL potassium.";
 RN Hum. Mol. Genet. 2:1349-1353(1993).
 RN [13]
 RP VARIANT POTASSIUM-AGGRAVATED MYOTONIA VAL-1160.
 RX PubMed=8058156;
 RA Ptacek L.J., Tawil R., Griggs R.C., Meola G., McManis P., Barohn R.J.,
 RT "Sodium channel mutations in acetazolamide-responsive myotonia
 RL congenita, paramyotonia congenita, and hyperkalemic periodic
 RN Neurology 44:1500-1503(1994).
 RN [14]
 RP VARIANT PARAMYOTONIA WITHOUT COLD PARALYSIS ILE-1293.
 RX MEDLINE=96154961; PubMed=8580427;
 RA Koch M.C., Baumbach K., George A.L., Ricker K.,
 RT "Paramyotonia congenita without paralysis on exposure to cold: a novel
 RL mutation in the SCN4A gene (Val1293Ile).";
 RN NeuroReport 6:2001-2004(1995).

RN [15]
 RP VARIANT POTASSIUM-AGGRAVATED MYOTONIA MET-445.
 RX PubMed=9392983;
 RA Rosenfeld J., Sloan-Brown K., George A.L. Jr.,
 RT "A novel muscle sodium channel mutation causes painful congenital
 RL myotonia.";
 RN Ann. Neurol. 42:811-814(1997).
 RN [16]
 RP VARIANT POTASSIUM-AGGRAVATED MYOTONIA MET-445.
 RX PubMed=10218481;
 RA Wang D.W., Vandecastee D., Ruben P.C., George A.L. Jr., Bennett P.B.,
 RT "Functional consequences of a domain 1/S6 segment sodium channel
 RL mutation associated with painful congenital myotonia.";
 RN FEBS Lett. 448:231-234(1999).
 RN [17]
 RP VARIANT HYPOKPP HIS-669.
 RX PubMed=10599760;
 RA Bulman D.E., Scoggan K.A., van Oene M.D., Nicollie M.W., Hahn A.F.,
 RT "A novel sodium channel mutation in a family with hypokalemic periodic
 RL paralysis.";
 RN Neurology 53:1932-1936(1999).
 RN [18]
 RP VARIANT HYPOKPP SER-1158.
 RX PubMed=10851391;
 RA Sugitara Y., Aoki T., Sugiyama Y., Hida C., Ogata M., Yamamoto T.,
 RT "Temperature-sensitive sodium channelopathy with heat-induced myotonia
 RL and cold-induced paralysis.";
 RN Neurology 54:2179-2181(2000).
 RN [19]
 RP VARIANT HYPOKPP GLY-672 AND HIS-672.
 RX PubMed=10944223;
 RA Jurkat-Rott K., Mitrovic N., Hang C., Kouzmine A., Ialzo P.,
 RT "Voltage-sensor sodium channel mutations cause hypokalemic periodic
 RL paralysis type 2 by enhanced inactivation and reduced current.";
 RN Proc. Natl. Acad. Sci. U.S.A. 97:9549-9554(2000).
 RN [20]
 RP VARIANT HYPOKPP SER-672.
 RX PubMed=11558801;
 RA Bendahhou S., Cummins T.R., Griggs R.C., Fu Y.H., Pracek L.J.,
 RT "Sodium channel inactivation defects are associated with
 RL acetazolamide-exacerbated hypokalemic periodic paralysis.";
 RN Ann. Neurol. 50:417-420(2001).
 RN [21]
 RP VARIANT HYPOKPP SER-672.
 RX PubMed=11591859;
 RA Davies N.P., Bunson L.H., Samuel M., Hanna M.G.,
 RT "Sodium channel gene mutations in hypokalemic periodic paralysis: an
 RL uncommon cause in the UK.";
 RN Neurology 57:1323-1325(2001).
 CC -1- FUNCTION: This protein mediates the voltage-dependent sodium ion
 CC permeability of excitable membranes. Assuming opened or closed
 CC conformations in response to the voltage difference across the
 CC membrane, the protein forms a sodium-selective channel through
 CC which Na⁺ ions may pass in accordance with their electrochemical
 CC gradient. This sodium channel may be present in both denervated
 CC and innervated skeletal muscle.
 CC -1- SUBUNIT: Muscle sodium channels contain an alpha subunit and a
 CC smaller beta subunit. Interacts with the PDZ domain of the
 CC syntrophin SNTN1, SNTB1 and SNTB2 (By similarity).
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
 CC -1- DOMAIN: The sequence contains 4 internal repeats, each with 5
 CC hydrophobic segments (S1, S2, S3, S5, S6) and one positively charged
 CC segment (S4). Segments S4 are probably the voltage-sensors and are
 CC characterized by a series of positively charged amino acids at
 CC every third position.
 CC -1- DISEASE: Defects in SCN4A are the cause of paramyotonia congenita
 CC of von Bultenbourg (PWC) [MIM:168300]. PWC is an autosomal dominant
 CC sodium channel disease characterized by myotonia, increased by
 CC exposure to cold, intermittent flaccid paresis, not necessarily

Query Match 100.0%; Score 28; DB 1; Length 1836;
 Best Local Similarity 100.0%; Pred. No. 5,1e-19;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MYLYFVFIIFGSEFTNLFGVIIDNF 28
 DB 1271 MYLYFVFIIFGSEFTNLFGVIIDNF 1298

RESULT 7
 ID CINA_RAT STANDARD; PRT; 1840 AA.
 AC P15350;
 DT 01-APR-1990 (Rel. 14, Last sequence update)
 DT 01-APR-1990 (Rel. 14, Last sequence update)
 DT 05-JUL-2004 (Rel. 44, Last annotation update)
 DE Sodium channel protein type IV alpha subunit (Voltage-gated sodium channel alpha subunit Nav1.4) (Sodium channel protein, skeletal muscle alpha-subunit) (Mu-1).
 GN Name=Scn4a;
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 OX NCBI_TaxId=10116;
 RN [1]
 RP MEDLINE=90148778; PubMed=2559760.
 RA Trimmer J.S., Cooperman S.S., Tomiko S.A., Zhou J., Crean S.M., Boyle M.B., Kallen R.G., Sheng Z., Barchi R.L., Sigworth F.J., Goodman R.H., Agnew W.S., Mandel G.,
 RT "Primary structure and functional expression of a mammalian skeletal muscle sodium channel."
 RT Neuron 33:33-49(1989).
 RL -1- FUNCTION: This protein mediates the voltage-dependent sodium ion permeability of excitable membranes. Assuming opened or closed conformations in response to the voltage difference across the membrane, the protein forms a sodium-selective channel through which Na+ ions may pass in accordance with their electrochemical gradient. This sodium channel may be present in both denervated and innervated skeletal muscle.
 CC -1- SUBUNIT: Muscle sodium channels contain an alpha subunit and a smaller beta subunit. Interacts with the PDZ domain of the synaptobrevin SNTA1, SNTB1 and SNTB2 (By similarity).
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
 CC -1- DOMAIN: The sequence contains 4 internal repeats, each with 5 hydrophobic segments (S1,S2,S3,S5,S6) and one positively charged segment (S4). Segments S4 are probably the voltage-sensors and are characterized by a series of positively charged amino acids at every third position.
 CC -1- SIMILARITY: Belongs to the sodium channel family.
 CC -1- SIMILARITY: Contains 1 IQ domain.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (See <http://www.ebi.ac.uk/announcements> or send an email to license@ebi.ac.uk).
 CC -----
 CC EMBL; M26643; AAA41682.1; -.
 DR PIR; JN0007; CHRTM1.
 DR HSSP; P04775; 1BYT.
 DR InterPro; IPR001682; Ca/Na_pore.
 DR InterPro; IPR002111; Cat_channel_TrpL.
 DR InterPro; IPR005821; Ion channel.
 DR InterPro; IPR000048; IQ region.
 DR InterPro; IPR005820; M-channel_nlg.
 DR InterPro; IPR001696; Na_channel.
 DR InterPro; IPR008052; Na_channel4.
 DR InterPro; IPR010526; Na_trans_assoc.
 DR Pfam; PF00520; Ion_trans_4.
 DR Pfam; PF00612; IQ_1.

DR Pfam; PF06512; Na_trans_assoc; 1.
 DR PRINTS; PRO0170; NACHANNEL.
 DR PRINTS; PRO1665; NACHANNEL4.
 DR PROSITE; PS50096; IQ_1.
 KW Glycoprotein; Ion transport; Ionic channel; Multigene family;
 KW Phosphorylation; Repeat; Sodium channel; Transmembrane;
 KW Voltage-gated channel.
 FT TRANSMEM 130 152 S1 of repeat I.
 FT TRANSMEM 156 179 S2 of repeat I.
 FT TRANSMEM 192 212 S3 of repeat I.
 FT TRANSMEM 214 233 S4 of repeat I.
 FT TRANSMEM 252 274 S5 of repeat I.
 FT TRANSMEM 417 444 S6 of repeat I.
 FT TRANSMEM 571 593 S1 of repeat II.
 FT TRANSMEM 663 685 S2 of repeat II.
 FT TRANSMEM 696 721 S3 of repeat II.
 FT TRANSMEM 722 737 S4 of repeat II.
 FT TRANSMEM 756 778 S5 of repeat II.
 FT TRANSMEM 832 859 S6 of repeat II.
 FT TRANSMEM 1084 1105 S1 of repeat III.
 FT TRANSMEM 1117 1140 S2 of repeat III.
 FT TRANSMEM 1149 1168 S3 of repeat III.
 FT TRANSMEM 1176 1195 S4 of repeat III.
 FT TRANSMEM 1215 1236 S5 of repeat III.
 FT TRANSMEM 1324 1351 S6 of repeat III.
 FT TRANSMEM 1405 1427 S1 of repeat IV.
 FT TRANSMEM 1437 1460 S2 of repeat IV.
 FT TRANSMEM 1468 1487 S3 of repeat IV.
 FT TRANSMEM 1502 1522 S4 of repeat IV.
 FT TRANSMEM 1535 1556 S5 of repeat IV.
 FT TRANSMEM 1627 1653 S6 of repeat IV.
 FT DOMAIN 1720 1749 IQ.
 FT CARBOHYD 288 288 N-linked (GlcNAc...)
 FT CARBOHYD 291 291 N-linked (GlcNAc...)
 FT CARBOHYD 297 297 N-linked (GlcNAc...)
 FT CARBOHYD 303 303 N-linked (GlcNAc...)
 FT CARBOHYD 309 309 N-linked (GlcNAc...)
 FT CARBOHYD 315 315 N-linked (GlcNAc...)
 FT CARBOHYD 327 327 N-linked (GlcNAc...)
 FT CARBOHYD 356 356 N-linked (GlcNAc...)
 FT CARBOHYD 502 502 N-linked (GlcNAc...)
 FT CARBOHYD 954 954 N-linked (GlcNAc...)
 FT CARBOHYD 1198 1198 N-linked (GlcNAc...)
 FT MOD_RES 56 56 Phosphoserine (by PKA)
 FT MOD_RES 251 251 Phosphoserine (by PKA)
 FT MOD_RES 1321 1321 Phosphoserine (by PKA)
 FT MOD_RES 1504 1504 Phosphoserine (by PKA)
 SQ SEQUENCE 1840 AA; 208865 MW; C5DC09D93DD9FAD6 CRC64;

Query Match 100.0%; Score 28; DB 1; Length 1840;
 Best Local Similarity 100.0%; Pred. No. 5,1e-19;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MYLYFVFIIFGSEFTNLFGVIIDNF 28
 DB 1264 MYLYFVFIIFGSEFTNLFGVIIDNF 1291

RESULT 8
 ID 070611 PRELIMINARY; PRT; 1840 AA.
 AC 070611;
 DT 01-AUG-1998 (TrEMBLrel. 07, Created)
 DT 01-AUG-1998 (TrEMBLrel. 07, Last sequence update)
 DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
 DE Rat skeletal muscle type 1 voltage-gated sodium channel (RSKM1) variant.
 GN Name=SCNA4;
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 OX NCBI_TaxId=10116;
 RN [1]

```

RP SEQUENCE FROM N.A.
RC STRAIN=Copenhagen; TISSUE=Prostate;
RX MEDLINE=98273645; PubMed=9613589;
RA Dias J.K.J., Stewart D., Fraser S.P., Black J.A., Dibb-Hajj S.,
RA Maxam S.G., Archer S.N., Djamgoz M.B.A.;
RT "Expression of skeletal muscle-type voltage-gated Na+ channel in rat
RT and human prostate cancer cell lines.";
RL FEBS Lett. 427:5-10(1998).
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
CC -1- SIMILARITY: Belongs to the sodium channel family.
DR EMBL; Y17153; CAA7659.1; -.
DR HSSP; P04775; IBBY.
DR GO; GO:0016021; C:Integral to membrane; IEA.
DR GO; GO:0005261; C:voltage-gated sodium channel complex; IEA.
DR GO; GO:0005248; F:cation channel activity; IEA.
DR GO; GO:0006814; F:voltage-gated sodium channel activity; IEA.
DR GO; GO:0006814; P:sodium ion transport; IEA.
DR GO; GO:0006814; P:sodium ion transport; IEA.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IQ_region.
DR InterPro; IPR005820; M_channel_nlg.
DR InterPro; IPR001696; Na_channel.
DR InterPro; IPR008052; Na_channel.
DR Pfam; PF00520; Ion_trans_4.
DR Pfam; PF00520; Ion_trans_4.
DR Pfam; PF06512; IQ_1.
DR PRINTS; PR00170; NACHANNEL.
DR PRINTS; PR01665; NACHANNEL4.
DR SMART; SM00015; IQ_1.
DR PROSITE; PS50096; IQ_1.
KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
KW Transport; Voltage-gated channel.
SQ SEQUENCE 1840 AA; 208823 MW; BIDFPA538E264B40 CRC64;

Query Match 100.0%; Score 28; DB 2; Length 1840;
Best Local Similarity 100.0%; Pred. No. 5,1e-19;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MYLYFVIFIFISFPTLNIFIGVIIDNF 28
Db 1264 MYLYFVIFIFISFPTLNIFIGVIIDNF 1291

RESULT 9
Q9ER60 PRELIMINARY; PRT; 1841 AA.
AC Q9ER60;
DT 01-MAR-2001 (TRENBLrel. 16, Created)
DT 01-MAR-2001 (TRENBLrel. 16, Last sequence update)
DE 01-MAR-2004 (TRENBLrel. 26, Last annotation update)
DE Voltage-gated sodium channel.
GN Name=Scn4a;
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxId=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=balb/c; TISSUE=Heart;
RX MEDLINE=21823196; PubMed=11834499;
RA Zimmer T., Bendorf K.;
RA "The mouse heart sodium channel (mH1): cloning and characterization of
RT alternatively spliced variants.";
RT Am. J. Physiol. Heart Circ. Physiol. 282:H1007-H1017(2002).
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
CC -1- SIMILARITY: Belongs to the sodium channel family.
DR EMBL; AJ278787; CAC17146.1; -.
DR HSSP; P04775; IBBY.
DR MGD; MGI:98250; Scn4a.
GO; GO:0016021; C:Integral to membrane; IEA.

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DR GO; GO:0005158; C:voltage-gated sodium channel complex; IEA.
DR GO; GO:0005261; F:cation channel activity; IEA.
DR GO; GO:0005248; F:voltage-gated sodium channel activity; IEA.
DR GO; GO:0006814; P:cation transport; IEA.
DR GO; GO:0006814; P:sodium ion transport; IEA.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR002111; Cat_channel_TripL.
DR InterPro; IPR005820; M_channel_nlg.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IQ_region.
DR InterPro; IPR001696; Na_channel.
DR InterPro; IPR008052; Na_channel.
DR Pfam; PF00520; Ion_trans_4.
DR Pfam; PF06512; IQ_1.
DR Pfam; PF06512; Na_trans_assoc; 1.
DR PRINTS; PR00170; NACHANNEL.
DR PRINTS; PR01665; NACHANNEL4.
DR SMART; SM00015; IQ_1.
DR PROSITE; PS50096; IQ_1.
KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
KW Transport; Voltage-gated channel.
SQ SEQUENCE 1841 AA; 208796 MW; 0766PDD3A9E0E55 CRC64;

Query Match 100.0%; Score 28; DB 2; Length 1841;
Best Local Similarity 100.0%; Pred. No. 5,1e-19;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MYLYFVIFIFISFPTLNIFIGVIIDNF 28
Db 1265 MYLYFVIFIFISFPTLNIFIGVIIDNF 1292

RESULT 10
CIN3_RAT STANDARD; PRT; 1951 AA.
ID CIN3_RAT
AC P08104;
DT 01-AUG-1988 (Rel. 08, Created)
DT 01-AUG-1988 (Rel. 08, Last sequence update)
DT 01-OCT-2004 (Rel. 45, Last annotation update)
DE Sodium channel subunit type III alpha subunit (Voltage-gated sodium
DE channel alpha subunit Nav1.3) (Sodium channel protein, brain III alpha
DE subunit) (Voltage-gated sodium channel subtype III).
GN Name=Scn3a;
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxId=10116;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Wistar;
RX MEDLINE=88137594; PubMed=249363;
RA Kayano T., Noda M., Flockerzi V., Takahashi H., Numa S.;
RA "Primary structure of rat brain sodium channel III deduced from the
RT cDNA sequence.";
RT FEBS Lett. 228:187-194(1988).
CC -1- FUNCTION: Mediates the voltage-dependent sodium ion permeability
CC of excitable membranes. Assuming opened or closed conformations in
CC response to the voltage difference across the membrane, the
CC protein forms a sodium-selective channel through which Na(+) ions
CC may pass in accordance with their electrochemical gradient.
CC -1- SUBUNIT: The sodium channel consists of a large polypeptide and 2-
CC 3 smaller ones. This sequence represents a large polypeptide.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
CC -1- DOMAIN: The sequence contains 4 internal repeats, each with 5
CC hydrophobic segments (S1,S2,S3,S5,S6) and one positively charged
CC segment (S4). Segments S4 are probably the voltage-sensors and are
CC characterized by a series of positively charged amino acids at
CC every third position.
CC -1- SIMILARITY: Belongs to the sodium channel family.
CC -1- SIMILARITY: Contains 1 IQ domain.
CC -----
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RP SEQUENCE FROM N.A.
RC TISSUE=Normal brain;
RA Ouchida M., Ohmori I.;
RL Submitted (DEC-2002) to the EMBL/GenBank/DBJ databases.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (by similarity).
DR HSP, P04775; IBY.
DR GO: GO:0016021; C: integral to membrane; IEA.
DR GO: GO:0001518; C: voltage-gated sodium channel complex; IEA.
DR GO: GO:0005261; F: cation channel activity; IEA.
DR GO: GO:0005248; F: voltage-gated sodium channel activity; IEA.
DR GO: GO:000612; P: cation transport; IEA.
DR GO: GO:000614; P: sodium ion transport; IEA.
DR InterPro: IPR01682; Ca/Na pore.
DR InterPro: IPR02111; Cat channel_TrpL.
DR InterPro: IPR005821; Ion trans.
DR InterPro: IPR000048; IO region.
DR InterPro: IPR005820; M-channel_nlg.
DR InterPro: IPR008051; Na_channel.
DR InterPro: IPR010526; Na_trans_assoc.
DR InterPro: IPR000100; Ribonuclease_P.
DR Pfam: PF00520; Ion trans; 4.
DR Pfam: PF00612; IO; 1.
DR Pfam: PF06512; Na trans assoc; 1.
DR PRINTS: PR00170; NACHANNEL.
DR PRINTS: PR01664; NACHANNEL.
DR SMART: SM00015; IO; 1.
DR PROSITE: PS00648; RIBONUCLEASE P; UNKNOWN 1.
KM Ion transport; Ionic channel; Sodium channel; Transmembrane;
KW Transport; Voltage-gated channel;
SQ SEQUENCE 1981 AA; 226201 MW; BID6946D6491B7AD CRC64;

Query Match 100.0%; Score 28; DB 2; Length 1981;
Best Local Similarity 100.0%; Pctd. No. 5.5e-19;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MYLFFVFIIFGSEFTLNFIVGIINP 28
DQ 1431 MYLFFVFIIFGSEFTLNFIVGIINP 1458

RESULT 13
CIN3_HUMAN
ID CIN3_HUMAN STANDARD; PRT; 2000 AA.
AC Q9NY46; Q9BZ83; Q9C066; Q9NYK2; Q9UPD1; Q9Y6P4;
DT 16-OCT-2001 (Rel. 40, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 01-OCT-2004 (Rel. 45, Last annotation update)
DE Sodium channel protein type III alpha subunit (Voltage-gated sodium channel alpha subunit Nav1.3) (Sodium channel protein, brain III alpha subunit) (Voltage-gated sodium channel subtype III).
GN Name=SCN3A; Synonyms=NA3;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_Taxid=9606;
RN [1]
RP SEQUENCE FROM N.A. (ISOFORM 2).
RC TISSUE=Brain;
RA Chen Y., Dale T.J., Romanos M.A., Whitaker W.R., Xie X., Clare J.J.;
RT "Cloning, distribution and functional analysis of the human brain type III sodium channel from human brain."
RL Submitted (DEC-1999) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A. (ISOFORM 3).
RA Jeong S.-Y., Goto J., Kanazawa I.;
RT "Cloning of cDNA for human voltage-gated sodium channel alpha subunit, SCN3A."
RL Submitted (JAN-2000) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A. (ISOFORMS 1; 2; 3 AND 4), AND VARIANT THR-606.

RA MEDLINE=21142400; PubMed=11245985; DOI=10.1016/S0378-1119(00)00594-1;
RC Kasai N., Fukushima K., Ueki Y., Prasad S., Nosakowski J.,
RA Sugata K.-I., Sugata A., Nishizaki K., Meyer N.C., Smith R.J.H.;
RT "Genomic structures of SCN2A and SCN3A - candidate genes for deafness at the DFNB16 locus."
RL Gene 264:113-122(2001).
RN [4]
RP SEQUENCE OF 1-1415 FROM N.A. (ISOFORMS 2 AND 4).
RC TISSUE=Brain;
RX MEDLINE=98251277; PubMed=9589372;
RA Lu C.M., Brown G.B.;
RT "Isolation of a human-brain sodium-channel gene encoding two isoforms of the subtype III alpha-subunit."
RL J. Mol. Neurosci. 10:67-70(1998).
RN [5]
RP SEQUENCE OF 1324-1413 FROM N.A.
RC TISSUE=Placenta;
RX MEDLINE=94211784; PubMed=8159690;
RA Malo M.S., Stivastava K., Andresen J.M., Chen X.N., Korenberg J.R.,
RT "Targeted gene walking by low stringency polymerase chain reaction: assignment of a putative human brain sodium channel gene (SCN3A) to chromosome 2q24-31."
RL Proc. Natl. Acad. Sci. U.S.A. 91:2975-2979(1994).
RN [6]
RP SEQUENCE OF 1669-1750 FROM N.A.
RC TISSUE=Kidney;
RA Tonkovich G.S., Kyle J.W.;
RT "Endogenous sodium current in HEK293 cells: increase in cell surface expression of endogenous currents by stable transfection of the Beta 1 subunit."
RL Submitted (FEB-2000) to the EMBL/GenBank/DBJ databases.
RN [7]
RP FUNCTION: Mediates the voltage-dependent sodium ion permeability of excitable membranes. Assuming opened or closed conformations in response to the voltage difference across the membrane, the protein forms a sodium-selective channel through which Na(+) ions may pass in accordance with their electrochemical gradient.
CC SUBUNIT: The sodium channel consists of a large polypeptide and 2-3 smaller ones. This sequence represents a large polypeptide.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
CC -1- ALTERNATIVE PRODUCTS:
CC Event=Alternative splicing; Named isoforms=4;
CC Comment=Exons 6A and 6N only differ by a single residue;
CC Name=1; Synonyms=6A-12+12b;
CC IsoId=Q9NY46-1; Sequence=Displayed;
CC Name=2; Synonyms=6A-12;
CC IsoId=Q9NY46-2; Sequence=VSP_001034;
CC Name=3; Synonyms=6N-12+12b;
CC IsoId=Q9NY46-3; Sequence=VSP_001033;
CC Name=4; Synonyms=6N-12;
CC IsoId=Q9NY46-4; Sequence=VSP_001033;
CC -1- DOMAIN: The sequence contains 4 internal repeats, each with 5 hydrophobic segments (S1,S2,S3,S5,S6) and one positively charged segment (S4). Segments S4 are probably the voltage-sensors and are characterized by a series of positively charged amino acids at every third position.
CC -1- SIMILARITY: Belongs to the sodium channel family.
CC -1- SIMILARITY: Contains 1 IO domain.
CC -----
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CC -----
DR EMBL: AJ251507; CAB5895.1; -;
DR EMBL: AF225987; AAK00219.1; -;
DR EMBL: AF330135; AAG53414.1; -;
DR EMBL: AF330118; AAG53414.1; JOINED.
DR EMBL: AF330119; AAG53414.1; JOINED.
DR EMBL: AF330120; AAG53414.1; JOINED.

FT	TRANSMEM	1297	1318	S4 of repeat III.
FT	TRANSMEM <td>1338</td> <td>1359</td> <td>S5 of repeat III.</td>	1338	1359	S5 of repeat III.
FT	TRANSMEM <td>1442</td> <td>1468</td> <td>S6 of repeat III.</td>	1442	1468	S6 of repeat III.
FT	TRANSMEM <td>1522</td> <td>1545</td> <td>S1 of repeat IV.</td>	1522	1545	S1 of repeat IV.
FT	TRANSMEM <td>1557</td> <td>1580</td> <td>S2 of repeat IV.</td>	1557	1580	S2 of repeat IV.
FT	TRANSMEM <td>1587</td> <td>1610</td> <td>S3 of repeat IV.</td>	1587	1610	S3 of repeat IV.
FT	TRANSMEM <td>1621</td> <td>1642</td> <td>S4 of repeat IV.</td>	1621	1642	S4 of repeat IV.
FT	TRANSMEM <td>1658</td> <td>1680</td> <td>S5 of repeat IV.</td>	1658	1680	S5 of repeat IV.
FT	TRANSMEM <td>1747</td> <td>1771</td> <td>S6 of repeat IV.</td>	1747	1771	S6 of repeat IV.
FT	DOMAIN <td>1900</td> <td>1929</td> <td>IQ.</td>	1900	1929	IQ.
FT	CARBOHYD <td>211</td> <td>211</td> <td>N-linked (GLCNAC. .)</td>	211	211	N-linked (GLCNAC. .)
FT	CARBOHYD <td>290</td> <td>290</td> <td>N-linked (GLCNAC. .)</td>	290	290	N-linked (GLCNAC. .)
FT	CARBOHYD <td>296</td> <td>296</td> <td>N-linked (GLCNAC. .)</td>	296	296	N-linked (GLCNAC. .)
FT	CARBOHYD <td>302</td> <td>302</td> <td>N-linked (GLCNAC. .)</td>	302	302	N-linked (GLCNAC. .)
FT	CARBOHYD <td>307</td> <td>307</td> <td>N-linked (GLCNAC. .)</td>	307	307	N-linked (GLCNAC. .)
FT	CARBOHYD <td>339</td> <td>339</td> <td>N-linked (GLCNAC. .)</td>	339	339	N-linked (GLCNAC. .)
FT	CARBOHYD <td>624</td> <td>624</td> <td>N-linked (GLCNAC. .)</td>	624	624	N-linked (GLCNAC. .)
FT	CARBOHYD <td>884</td> <td>884</td> <td>N-linked (GLCNAC. .)</td>	884	884	N-linked (GLCNAC. .)
FT	CARBOHYD <td>1051</td> <td>1051</td> <td>N-linked (GLCNAC. .)</td>	1051	1051	N-linked (GLCNAC. .)
FT	CARBOHYD <td>1068</td> <td>1068</td> <td>N-linked (GLCNAC. .)</td>	1068	1068	N-linked (GLCNAC. .)
FT	CARBOHYD <td>1134</td> <td>1134</td> <td>N-linked (GLCNAC. .)</td>	1134	1134	N-linked (GLCNAC. .)
FT	CARBOHYD <td>1366</td> <td>1366</td> <td>N-linked (GLCNAC. .)</td>	1366	1366	N-linked (GLCNAC. .)
FT	CARBOHYD <td>1380</td> <td>1380</td> <td>N-linked (GLCNAC. .)</td>	1380	1380	N-linked (GLCNAC. .)
FT	VARSPPLIC <td>208</td> <td>208</td> <td>S -> D (in isoform 3 and isoform 4).</td>	208	208	S -> D (in isoform 3 and isoform 4).
FT	VARSPPLIC <td>625</td> <td>673</td> <td>/FTId=VSP_001033.</td>	625	673	/FTId=VSP_001033.
FT	VARSPPLIC <td>625</td> <td>673</td> <td>Missing (in isoform 2 and isoform 4).</td>	625	673	Missing (in isoform 2 and isoform 4).

Query Match 100.0%; Score 28; DB 1; Length 2000;
Beat Local Similarity 100.0%; Pred. No. 5..5e-19;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0

Qy 1 MLYLTVFTITFGSGFPLTLNLTFTGIYIINDF 28
Db 1444 MLYLTVFTITFGSGFPLTLNLTFTGIYIINDF 1471

RESULT 14
CIN2_HUMAN STANDARD; PRT; 2005 AA.
ID CIN2_HUMAN
AC Q99250; Q14472; Q9BZC9; Q9BZD0;
DT 01-JUN-1994 (Rel. 29, Created)
DT 26-FEB-2003 (Rel. 41, Last sequence update)
DT 05-JUL-2004 (Rel. 44, Last annotation update)
DE Sodium channel protein type II alpha subunit (Voltage-gated sodium channel alpha subunit Nav1.2) (Sodium channel protein, brain II alpha subunit) (HBSG II).
DN Name=SCN2A; Synonyms=SCN2A2, NAC2;
GN Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A. (ISOFORM 1).
RC TISSUE=Brain;
RX MEDLINE=92300418; PubMed=1325650;
RA Ahmed C.M., Ware D.H., Lee S.C., Patten C.D., Ferrer-Montiel A.V., Schindler A.F., McPherson J.D., Wagner-McPherson C.B., Wasmuth J.J., Evans G.A., Montal M.;
RT "Primary structure, chromosomal localization, and functional expression of a voltage-gated sodium channel from human brain."; Proc. Natl. Acad. Sci. U.S.A. 89:8220-8224(1992).
RL [2]
RP SEQUENCE FROM N.A. (ISOFORMS 1 AND 2).
RX MEDLINE=21142400; PubMed=11245985; DOI=10.1016/S0378-1119(00)00594-1;
RA Kasai N., Fukushima K., Ueki Y., Prasad S., Nozakiowski J., Sugata K.-I., Sugata A., Nishizaki K., Meyer N.C., Smith R.J.H.;
RT "Genomic structures of SCN2A and SCN3A - candidate genes for deafness at the D2NA16 locus.";
RL Gene 264:113-122(2001).
RN [3]
RP SEQUENCE OF 1-89 FROM N.A.
RT Lu C.-M., Eichelberger U.S., Beckman M.L., Schade S.D., Brown G.B.;
RT "Isolation of the 5'-flanking region for human brain sodium channel

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RT subtype II alpha-Subunit (SCN2A).";
RL Submitted (APR-1998) to the EMBL/GenBank/DBJ databases.
RP [4]
RX SEQUENCE OF 1702-2005 FROM N.A.
RC TISSUE=Brain.
RX MEDLINE=92275082; Pubmed=1317301;
RA Lu C.-M., Han U., Rado T.A., Brown G.B.;
RT "Differential expression of two sodium channel subtypes in human
RL brain.";
RT FEBS Lett. 303:53-58(1992).
RN [5]
RP SEQUENCE OF 1702-1772 FROM N.A.
RX MEDLINE=9110524; Pubmed=1846440;
RA Han U., Lu C.-M., Brown G.B., Rado T.A.;
RT "Direct amplification of a single dissected chromosomal segment by
RT polymerase chain reaction: a human brain sodium channel gene is on
RL Chromosome 2q22-q23.";
RL Proc. Natl. Acad. Sci. U.S.A. 88:335-339(1991).
CC -1- FUNCTION: Mediates the voltage-dependent sodium ion permeability
CC of excitable membranes. Assuming opened or closed conformations in
CC response to the voltage difference across the membrane, the
CC protein forms a sodium-selective channel through which Na(+) ions
CC may pass in accordance with their electrochemical gradient.
CC -1- SUBUNIT: The sodium channel consists of a large polypeptide and 2-
CC 3 smaller ones. This sequence represents a large polypeptide.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
CC -1- ALTERNATIVE PRODUCTS:
CC Event=Alternative splicing; Named isoforms=2;
CC Name=1; Synonyms=Adult, 6A;
CC IsoId=Q99250-1; Sequence=Neonatal, 6N;
CC Name=2; Synonyms=Neonatal, 6N;
CC IsoId=Q99250-2; Sequence=VSP_001032;
CC -1- DOMAIN: The sequence contains 4 internal repeats, each with 5
CC hydrophobic segments (S1,S2,S3,S5,S6) and one positively charged
CC segment (S4). Segments S4 are probably the voltage-sensors and are
CC characterized by a series of positively charged amino acids at
CC every third position.
CC -1- SIMILARITY: Belongs to the sodium channel family.
CC -1- SIMILARITY: Contains 1 IQ domain.
CC
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CC or send an email to license@isb-sib.ch).
CC
CC -----
DR EMBL; M94055; AAA18895.1; -
DR EMBL; AF059683; AAC14574.1; -
DR EMBL; AF327246; AAG53413.1; -
DR EMBL; AF327226; AAG53413.1; JOINED.
DR EMBL; AF327227; AAG53413.1; JOINED.
DR EMBL; AF327228; AAG53413.1; JOINED.
DR EMBL; AF327229; AAG53413.1; JOINED.
DR EMBL; AF327230; AAG53413.1; JOINED.
DR EMBL; AF327231; AAG53413.1; JOINED.
DR EMBL; AF327232; AAG53413.1; JOINED.
DR EMBL; AF327233; AAG53413.1; JOINED.
DR EMBL; AF327234; AAG53413.1; JOINED.
DR EMBL; AF327235; AAG53413.1; JOINED.
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DR EMBL; AF327237; AAG53413.1; JOINED.
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DR EMBL; AF327239; AAG53413.1; JOINED.
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DR EMBL; AF327242; AAG53413.1; JOINED.
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DR EMBL; AF327246; AAG53412.1; -
DR EMBL; AF327226; AAG53412.1; JOINED.
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DR EMBL; AF327233; AAG53412.1; JOINED.
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DR EMBL; AF327235; AAG53412.1; JOINED.
DR EMBL; AF327236; AAG53412.1; JOINED.
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DR EMBL; AF327239; AAG53412.1; JOINED.
DR EMBL; AF327240; AAG53412.1; JOINED.
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DR EMBL; AF327243; AAG53412.1; JOINED.
DR EMBL; AF327244; AAG53412.1; JOINED.
DR EMBL; AF327245; AAG53412.1; JOINED.
DR EMBL; M91804; -; NOT ANNOTATED_CDS.
DR EMBL; M55662; AAB65854.2; -
DR HSSP; P04775; 1BYX.
DR Gene: HGNC:10588; SCN2A2.
DR MIM; 601219; -
DR GO; GO:0005887; C:integral to plasma membrane; TAS.
DR GO; GO:0005248; F:voltage-gated sodium channel activity; TAS.
DR GO; GO:0006814; P:sodium ion transport; TAS.
DR InterPro; IPR001682; Ca/Na.pore.
DR InterPro; IPR002111; Cat_channel_TripL.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IQ_region.
DR InterPro; IPR005820; M+channel_nlg.
DR InterPro; IPR001696; Na_channel.
DR InterPro; IPR010526; Na_trans_assoc.
DR Pfam; PF00520; Ion_trans_4.
DR Pfam; PF00612; IQ_1.
DR Pfam; PF06512; Na_trans_assoc; 1.
DR PRINTS; PR00170; NACHANNEL.
DR PROSITE; PSS0096; IQ_1.
DR KX Alternative splicing; Glycoprotein; Ion transport; Ionic channel;
DR KX Multigene family; Repeat; Sodium channel; Transmembrane;
DR KW Voltage-gated channel.
FT REPEAT 111 456
FT REPEAT 741 1013
FT REPEAT 1190 1504
FT REPEAT 1513 1811
FT TRANSMEM 125 148
FT TRANSMEM 157 176
FT TRANSMEM 190 208
FT TRANSMEM 215 234
FT TRANSMEM 251 274
FT TRANSMEM 402 427
FT TRANSMEM 754 778
FT TRANSMEM 790 813
FT TRANSMEM 822 841
FT TRANSMEM 848 867
FT TRANSMEM 884 904
FT TRANSMEM 958 983
FT TRANSMEM 1204 1227
FT TRANSMEM 1241 1266
FT TRANSMEM 1273 1294
FT TRANSMEM 1299 1320
FT TRANSMEM 1340 1367
FT TRANSMEM 1447 1473
FT TRANSMEM 1527 1550
FT TRANSMEM 1562 1585
FT TRANSMEM 1592 1615
FT TRANSMEM 1626 1647
FT TRANSMEM 1663 1685
FT TRANSMEM 1752 1776
FT DOMAIN 1905 1934
FT CARBOHYD 212 212

```

N-linked (GlcNAc. . .) (Potential).


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FT CARBOHYD 285 285 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 291 291 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 297 297 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 303 303 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 308 308 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 340 340 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 604 604 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 624 624 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 883 883 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 1055 1055 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 1072 1072 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 1136 1136 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 1368 1368 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 1382 1382 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 1393 1393 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 1778 1778 N-linked (GlcNAc... ) (Potential)
FT VARSPLIC 209 209 D -> N (in isoform 2).
FT CONFLICT 524 524 /FTId=VSP_001032.
R -> L (in Ref. 1).

Query Match 100.0%; Score 28; DB 1; Length 2005;
Best Local Similarity 100.0%; Pred. No. 5.5e-19;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 MLYVFIIFIGSFPTLNFIVGIIDNF 28
Db 1449 MLYVFIIFIGSFPTLNFIVGIIDNF 1476

RESULT 15
ID CIN2 RAT STANDARD; PRT; 2005 AA.
AC P04775;
DT 13-AUG-1987 (Rel. 05, Created)
DT 13-AUG-1987 (Rel. 05, Last sequence update)
DT 01-OCT-2004 (Rel. 45, Last annotation update)
DE Sodium channel protein type II alpha subunit (Voltage-gated sodium
channel alpha subunit Nav1.2) (Sodium channel protein, brain II alpha
subunit).
GN Name=Scn2a;
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OC NCBI_TaxId=10116;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=86146901; PubMed=3754035;
RA Node M., Ikeda T., Kayano T., Suzuki H., Takeshima H., Kurasaki M.,
RA Takahashi H., Numa S.;
RT "Existence of distinct sodium channel messenger RNAs in rat brain.";
RL Nature 320:186-192(1986).
CC -1- FUNCTION: Mediates the voltage-dependent sodium ion permeability
of excitable membranes. Assuming opened or closed conformations in
response to the voltage difference across the membrane, the
protein forms a sodium-selective channel through which Na(+) ions
may pass in accordance with their electrochemical gradient.
CC -1- SUBUNIT: The sodium channel consists of a large polypeptide and 2-
3 smaller ones. This sequence represents a large polypeptide.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
CC -1- DOMAIN: The sequence contains 4 internal repeats, each with 5
hydrophobic segments (S1,S2,S3,S5,S6) and one positively charged
segment (S4). Segments S4 are probably the voltage-sensors and are
characterized by a series of positively charged amino acids at
every third position.
CC -1- SIMILARITY: Belongs to the sodium channel family.
CC -1- SIMILARITY: Contains 1 IQ domain.
-----
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CC or send an email to license@isb-sib.ch).
CC EMBL: X03639; CAA27287.1; -
DR PDB; 1BY7; NMR; A=1474-1526.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR002111; Cat_channel_TrypL.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IQ_region.
DR InterPro; IPR005820; M-channel_nlg.
DR InterPro; IPR001696; Na_channel.
DR InterPro; IPR010526; Na_trans_assoc.
DR Pfam; PF00520; Ion_trans; 4.
DR Pfam; PF06512; IQ; 1.
DR Pfam; PF06512; Na_trans_assoc; 1.
DR PRINTS; PR00170; NACHANNEL.
DR PROSITE; PS50096; IQ; 1.
KW 3D-structure; Glycoprotein; Ion transport; Ionic channel;
KW Multigene family; Repeat; Sodium channel; Transmembrane;
KW Voltage-gated channel.
FT REPEAT 111 456 I.
FT REPEAT 741 1013 II.
FT REPEAT 1190 1504 III.
FT REPEAT 1513 1811 IV.
FT TRANSMEM 125 148 S1 of repeat I.
FT TRANSMEM 157 176 S2 of repeat I.
FT TRANSMEM 190 208 S3 of repeat I.
FT TRANSMEM 215 234 S4 of repeat I.
FT TRANSMEM 251 274 S5 of repeat I.
FT TRANSMEM 402 427 S6 of repeat I.
FT TRANSMEM 754 778 S1 of repeat II.
FT TRANSMEM 813 833 S2 of repeat II.
FT TRANSMEM 822 841 S3 of repeat II.
FT TRANSMEM 848 867 S4 of repeat II.
FT TRANSMEM 884 904 S5 of repeat II.
FT TRANSMEM 958 983 S6 of repeat II.
FT TRANSMEM 1204 1227 S1 of repeat III.
FT TRANSMEM 1241 1266 S2 of repeat III.
FT TRANSMEM 1273 1294 S3 of repeat III.
FT TRANSMEM 1299 1320 S4 of repeat III.
FT TRANSMEM 1340 1367 S5 of repeat III.
FT TRANSMEM 1447 1473 S6 of repeat III.
FT TRANSMEM 1527 1550 S1 of repeat IV.
FT TRANSMEM 1562 1585 S2 of repeat IV.
FT TRANSMEM 1592 1615 S3 of repeat IV.
FT TRANSMEM 1626 1647 S4 of repeat IV.
FT TRANSMEM 1663 1685 S5 of repeat IV.
FT TRANSMEM 1752 1776 S6 of repeat IV.
FT DOMAIN 1905 1934 IQ.
FT CARBOHYD 212 212 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 285 285 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 291 291 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 297 297 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 303 303 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 308 308 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 340 340 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 360 360 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 604 604 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 624 624 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 883 883 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 1055 1055 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 1072 1072 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 1136 1136 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 1368 1368 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 1382 1382 N-linked (GlcNAc... ) (Potential)
FT CARBOHYD 1393 1393 N-linked (GlcNAc... ) (Potential)
SQ SEQUENCE 2005 AA; 227872 MW; 861B583D79F8324 CRC64;

Query Match 100.0%; Score 28; DB 1; Length 2005;
Best Local Similarity 100.0%; Pred. No. 5.5e-19;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 MLYVFIIFIGSFPTLNFIVGIIDNF 28
Db 1449 MLYVFIIFIGSFPTLNFIVGIIDNF 1476

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RESULT 16
09YGN7 PRELIMINARY; PRT; 2007 AA.
AC 09YGN7;
DT 01-MAY-1999 (TREMBlrel. 10, Created)
DT 01-MAY-1999 (TREMBlrel. 10, Last sequence update)
DT 01-MAR-2004 (TREMBlrel. 26, Last annotation update)
DE Voltage-dependent sodium channel.
OS Cynops pyrrhogaster (Japanese common newt).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Amphibia; Batrachia; Caudata; Salamandroidea; Salamandridae; Cynops.
OX NCBI_TaxId=8330;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Retina;
RA Hirota K., Kaneko Y., Matsumoto G., Hanyu Y.;
RL Submitted (JAN-1999) to the EMBL/GenBank/DBJ databases.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
CC EMBL; AF123593; AAD17315.1; -.
DR HSBP; P04775; IBY.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0003158; C:voltage-gated sodium channel complex; IEA.
DR GO; GO:0005261; F:cation channel activity; IEA.
DR GO; GO:0005248; F:voltage-gated sodium channel activity; IEA.
DR GO; GO:0006812; P:cation transport; IEA.
DR GO; GO:0006814; P:sodium ion transport; IEA.
DR InterPro: IPR001682; Ca/Na pore.
DR InterPro: IPR002111; Cat_channel_TrpL.
DR InterPro: IPR005821; Ion_trans.
DR InterPro: IPR000048; IQ_region.
DR InterPro: IPR005820; M_channel_nlg.
DR InterPro: IPR001696; Na_channel.
DR InterPro: IPR010526; Na_trans_assoc.
DR Pfam; PF00520; Ion_trans_4.
DR Pfam; PF00612; IQ_1.
DR Pfam; PF06512; Na_trans_assoc_1.
DR PRINTS; PR00170; NACHANNEL.
DR SMART; SM00015; IQ_1.
KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
KW Transport; Voltage-gated channel.
SQ SEQUENCE 2007 AA; 228398 MW; 0135B9B9EC9C294C9 CRC64;

Query Match 100.0%; Score 28; DB 2; Length 2007;
Best Local Similarity 100.0%; Pred. No. 5.5e-19;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 MLYLPIFIIPSGFPTLNPIGVIIIDNF 28
Db 1450 MLYLPIFIIPSGFPTLNPIGVIIIDNF 1477

RESULT 17
CINI_HUMAN STANDARD; PRT; 2009 AA.
AC P35498; Q16172; Q96LA3; Q9CC08;
DT 01-JUN-1994 (Rel. 29, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 05-JUL-2004 (Rel. 44, Last annotation update)
DE Sodium channel protein type I alpha subunit (Voltage-gated sodium
DE channel alpha subunit Nav1.1) (Sodium channel protein, Brain I alpha
DE subunit).
GN Name=SCN1A, Synonyms=SCN1, NAC1;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxId=9606;
RN [1]
RP SEQUENCE FROM N.A. (ISOFORM 1), AND VARIANTS GEFS+2 MET-875 AND
RP HIS-1648.
RX MEDLINE=20206553; PubMed=10742094;

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RA Escayz A., MacDonald B.T., Meisler M.H., Baulac S., Huberfeld G.,
RA An-Gourfinkel I., Brice A., LeGuern E., Moulard B., Chaigne D.,
RA Buresti C., Malafosse A.;
RT "Mutations of SCN1A, encoding a neuronal sodium channel, in two
RT families with GEFS+2."
RL Nat. Genet. 24:343-345(2000).
RN [2]
RP SEQUENCE FROM N.A. (ISOFORM 2).
RA Jeong S.-Y., Goto J., Kanazawa I.;
RT "Cloning of cDNA for human voltage-gated sodium channel alpha subunit,
RT SCN1A."
RL Submitted (JAN-2000) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A. (ISOFORM 2).
RC TISSUE=Brain;
RA Sugawara T., Mazaki E.M., Yamakawa K.;
RT "Homo sapiens neuronal voltage-gated sodium channel type I (Nav1.1)
RT mRNA."
RL Submitted (JUL-2001) to the EMBL/GenBank/DBJ databases.
RN [4]
RP SEQUENCE FROM N.A. (ISOFORMS 1 AND 2), AND VARIANT THR-1067.
RA Ouchida M., Ohmori I.;
RT "Isoforms of human sodium channel SCN1A gene."
RL Submitted (OCT-2002) to the EMBL/GenBank/DBJ databases.
RN [5]
RP SEQUENCE OF 1335-1428 FROM N.A.
RX MEDLINE=94340991; PubMed=8062593;
RA Malo M.S., Blanchard B.J., Andresen J.M., Srivastava K., Chen X.N.,
RA Li X., Jabs E.W., Korenberg J.R., Ingram V.M.;
RT "Localization of a putative human brain sodium channel gene (SCN1A) to
RT chromosome band 2q24."
RL Cytogenet. Cell Genet. 67:178-186(1994).
RN [6]
RP SEQUENCE OF 1518-1940 FROM N.A.
RC TISSUE=Brain;
RX MEDLINE=92275082; PubMed=1317301;
RA Lu C.-M., Han J., Rado T.A., Brown G.B.;
RT "Differential expression of two sodium channel subtypes in human
RT brain."
RL FEBS Lett. 303:53-58(1992).
RN [7]
RP VARIANTS GEFS+2 VAL-188; LEU-1353 AND MET-1656.
RX MEDLINE=21152274; PubMed=11254444;
RA Wallace R.H., Scheffer I.E., Barnett S., Richards M., Dibbens L.,
RA Desai R.R., Lerman-Sagie T., Lev D., Mazariu A., Brand N.,
RA Ben-Zeev B., Golkman I., Singh R., Kremidioti G., Gardner A.,
RA Suberland G.R., George A.L. Jr., Mulley J.C., Berkovic S.F.;
RT "Neuronal sodium-channel alpha1-subunit mutations in generalized
RT epilepsy with febrile seizures plus."
RL Am. J. Hum. Genet. 68:859-865(2001).
RN [8]
RP VARIANT GEFS+2 ARG-1204.
RX MEDLINE=2152275; PubMed=11254445;
RA Escayz A., Heils A., MacDonald B.T., Haug K., Sander T., Meisler M.H.;
RT "A novel SCN1A mutation associated with generalized epilepsy with
RT febrile seizures plus -- and prevalence of variants in patients with
RT epilepsy."
RL Am. J. Hum. Genet. 68:866-873(2001).
RN [9]
RP VARIANT SMEI PHE-986.
RX MEDLINE=21257503; PubMed=11359211;
RA Claes L., Del-Favero J., Ceulemans B., Lagae L., Van Broeckhoven C.,
RA De Jonghe P.;
RT "de novo mutations in the sodium-channel gene SCN1A cause severe
RT myoclonic epilepsy of infancy."
RL Am. J. Hum. Genet. 68:1327-1332(2001).
RN [10]
RP VARIANT GEFS+2 THR-1270.
RX MEDLINE=21630138; PubMed=11756608;
RA Abou-Khalil B., Ge Q., Desai R., Ryther R., Bazyk A., Bailey R.,
RA Haines J.L., Sutcliffe J.S., George A.L. Jr.;
RT "Partial and generalized epilepsy with febrile seizures plus and a
RT novel SCN1A mutation."

```

RL Neurology 57:2265-2272(2001).

CC -1- FUNCTION: Mediates the voltage-dependent sodium ion permeability

CC of excitable membranes. Assuming opened or closed conformations in

CC response to the voltage difference across the membrane, the

CC protein forms a sodium-selective channel through which Na(+) ions

CC may pass in accordance with their electrochemical gradient.

CC -1- SUBUNIT: The sodium channel consists of a large polypeptide and 2-

CC 3 smaller ones. This sequence represents a large polypeptide.

CC -1- SUBCELLULAR LOCATION: Integral membrane protein.

CC -1- ALTERNATIVE PRODUCTS: Integral membrane protein.

CC Event=Alternative splicing; Named isoforms=2;

CC Name=1;

CC IsoId=P35498-1; Sequence=Dieplayed;

CC Name=2;

CC IsoId=P35498-2; Sequence=VSP_001031;

CC Note=No experimental confirmation available;

CC -1- DOMAIN: The sequence contains 4 internal repeats, each with 5

CC hydrophobic segments (S1,S2,S3,S5,S6) and one positively charged

CC segment (S4). Segments S4 are probably the voltage-sensors and are

CC characterized by a series of positively charged amino acids at

CC every third position.

CC -1- DISEASE: Defects in SCN1A are the cause of generalized epilepsy

CC with febrile seizures plus type 2 (GFS-2) [MIM:604233]. This

CC autosomal dominant disorder is characterized by febrile seizures

CC in children and afebrile seizures in adults. Penetrance is

CC incomplete and a large intrafamilial variability of the phenotype

CC is observed.

CC -1- DISEASE: Defects in SCN1A are a cause of severe myoclonic epilepsy

CC in infancy (SMEI) [MIM:607208], a severe form of generalized

CC epilepsy with febrile seizures. SMEI is a rare disorder

CC characterized by normal development before onset, seizures

CC beginning in the first year of life in the form of generalized or

CC unilateral febrile clonic seizures, secondary appearance of

CC myoclonic seizures, and occasionally partial seizures. It is

CC associated with ataxia, slowed psychomotor development, and mental

CC decline.

CC -1- SIMILARITY: Belongs to the sodium channel family.

CC -1- SIMILARITY: Contains 1 IQ domain.

CC -----

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CC between the Swiss Institute of Bioinformatics and the EMBL outstation -

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CC -----

DR EMBL; AF225985; AAK00217.1; -

DR EMBL; AY043484; AAK95360.1; -

DR EMBL; AB093548; BAC21101.1; -

DR EMBL; AB093549; BAC21102.1; -

DR EMBL; S71446; AAB31605.1; -

DR EMBL; X65362; CAA46439.1; -

DR EMBL; M91803; -; NOT_ANNOTATED_CDS.

DR PIR; I52964; I52964.

DR PIR; S29184; S29184.

DR HSSP; P04775; 1BRY.

DR Genew; HGNC:10585; SCN1A.

DR MIM; 182389; -

DR MIM; 604233; -

DR MIM; 607208; -

DR GO; GO:0016021; C:integral to membrane; NAS.

DR GO; GO:0005248; F:voltage-gated sodium channel activity; NAS.

DR GO; GO:0006814; F:sodium ion transport; NAS.

DR InterPro; IPR001682; Ca/Na_pore.

DR InterPro; IPR002111; Cat_channel_Tryp.

DR InterPro; IPR005821; Ion_trans.

DR InterPro; IPR000048; IQ_region.

DR InterPro; IPR005820; M-channel_nlg.

DR InterPro; IPR001696; Na_channel1.

DR InterPro; IPR008051; Na_channel1.

DR InterPro; IPR010526; Na_trans_assoc.

DR Pfam; PF00520; Ion_trans; 4.

DR Pfam; PF00612; IQ; 1.

DR Pfam; PF06512; Na_trans_assoc; 1.

DR PRINTS; PR00170; NACHANNEL.

DR PRINTS; PR01664; NACHANNEL.

DR PROSITE; PS50096; IQ; FALSE_NEG.

DR Alternative splicing; Disease mutation; Epilepsy; Glycoprotein;

DR Ion transport; Ionic channel; Multigene family; Polymorphism; Repeat;

DR Sodium channel; Transmembrane; Voltage-gated channel.

FT REPEAT 110 454

FT REPEAT 750 1022

FT REPEAT 1200 1514

FT REPEAT 1523 1821

FT TRANSMEM 124 147

FT TRANSMEM 156 175

FT TRANSMEM 189 207

FT TRANSMEM 214 233

FT TRANSMEM 230 273

FT TRANSMEM 400 425

FT TRANSMEM 763 787

FT TRANSMEM 799 822

FT TRANSMEM 831 850

FT TRANSMEM 857 876

FT TRANSMEM 893 913

FT TRANSMEM 967 992

FT TRANSMEM 1214 1237

FT TRANSMEM 1251 1276

FT TRANSMEM 1283 1304

FT TRANSMEM 1309 1330

FT TRANSMEM 1350 1377

FT TRANSMEM 1457 1483

FT TRANSMEM 1537 1560

FT TRANSMEM 1572 1595

FT TRANSMEM 1602 1625

FT TRANSMEM 1636 1657

FT TRANSMEM 1673 1695

FT TRANSMEM 1762 1786

FT CARBOHYD 211 211

Query Match 100.0%; Score 28; DB 1; Length 2009;

Best Local Similarity 100.0%; Pred. No. 5.5e-19;

Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 MYLVFVPIIFGSPFTLNLFGIVITDNF 28

Db 1459 MYLVFVPIIFGSPFTLNLFGIVITDNF 1486

RESULT 18

CINI RAT STANDARD; PRT; 2009 AA.

AC P04774;

DT 13-AUG-1987 (Rel. 05, Created)

DT 13-AUG-1987 (Rel. 05, Last sequence update)

DT 05-JUL-2004 (Rel. 44, Last annotation update)

DE Sodium channel protein type I alpha subunit (Voltage-gated sodium

DE channel alpha subunit Nav1.1) (Sodium channel protein, brain I alpha

DE subunit).

GN Name=Scn1a;

OS Rattus norvegicus (Rat).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.

OX NCBI_TaxID=10116;

RN [1]

RN SEQUENCE FROM N.A.

RX MEDLINE=86146901; PubMed=3754035;

RA Noda M., Ikeda T., Kayano T., Suzuki H., Takeshima H., Kuraaki M.,

RA "Takeshima H., Numa S.;

RT Nature 320:188-192(1986).

RL [2]

RN SEQUENCE FROM N.A.

RX MEDLINE=87311395; PubMed=2442385;

RA Noda M., Numa S.;

RT "Structure and function of sodium channel.";
 CC J. Receptor. Res. 7:467-497(1987).
 CC -1- FUNCTION: Mediates the voltage-dependent sodium ion permeability
 CC of excitable membranes. Assuming opened or closed conformations in
 CC response to the voltage difference across the membrane, the
 CC protein forms a sodium-selective channel through which Na(+) ions
 CC may pass in accordance with their electrochemical gradient.
 CC -1- SUBUNIT: The sodium channel consists of a large polypeptide and 2-
 CC 3 smaller ones. This sequence represents a large polypeptide.
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
 CC -1- DOMAIN: The sequence contains 4 internal repeats, each with 5
 CC hydrophobic segments (S1,S2,S3,S5,S6) and one positively charged
 CC segment (S4). Segments S4 are probably the voltage-sensors and are
 CC characterized by a series of positively charged amino acids at
 CC every third position.
 CC -1- SIMILARITY: Belongs to the sodium channel family.
 CC -1- SIMILARITY: Contains 1 IQ domain.
 CC -----
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 CC or send an email to license@sib-sib.ch).
 CC -----
 DR EMBL; X03638; CAA27286.1; -;
 DR EMBL; M22253; AAA79965.1; -;
 DR PIR; A25019; A25019.
 DR HSSP; P04775; IBY.
 DR RGD; 69364; Scn1a.
 DR InterPro; IPR001682; Ca/Na pore.
 DR InterPro; IPR002111; Cat channel_TrpL.
 DR InterPro; IPR005821; Ion trans.
 DR InterPro; IPR000048; IQ region.
 DR InterPro; IPR005820; M-channel_nlg.
 DR InterPro; IPR001696; Na channel_nlg.
 DR InterPro; IPR008051; Na channel.
 DR InterPro; IPR010526; Na trans_assoc.
 DR Pfam; PF00520; Ion trans; 4.
 DR Pfam; PF00612; IQ; 1.
 DR Pfam; PF06512; Na trans_assoc; 1.
 DR PRINTS; PR00170; NACHANNEL.
 DR PRINTS; PR01664; NACHANNEL.
 DR PROSITE; PS50096; IQ; FALSE_NEG.
 KW Glycoprotein; Ion transport; Ionic channel; Multigene family; Repeat;
 KW Sodium channel; Transmembrane; Voltage-gated channel.
 FT REPEAT 110 454
 FT REPEAT 750 1022
 FT REPEAT 1200 1514
 FT REPEAT 1523 1821
 FT TRANSMEM 124 147
 FT TRANSMEM 156 175
 FT TRANSMEM 189 207
 FT TRANSMEM 214 233
 FT TRANSMEM 250 273
 FT TRANSMEM 400 425
 FT TRANSMEM 763 787
 FT TRANSMEM 799 822
 FT TRANSMEM 831 850
 FT TRANSMEM 857 876
 FT TRANSMEM 893 913
 FT TRANSMEM 967 992
 FT TRANSMEM 1214 1237
 FT TRANSMEM 1251 1276
 FT TRANSMEM 1283 1304
 FT TRANSMEM 1309 1330
 FT TRANSMEM 1350 1377
 FT TRANSMEM 1457 1483
 FT TRANSMEM 1537 1560
 FT TRANSMEM 1572 1595
 FT TRANSMEM 1602 1625
 FT TRANSMEM 1636 1657
 S4 of repeat IV.

FT TRANSMEM 1673 1695 S5 of repeat IV.
 FT TRANSMEM 1762 1786 S6 of repeat IV.
 FT CARBOHYD 211 211 N-linked (GlcNAc...)
 FT CARBOHYD 284 284 N-linked (GlcNAc...)
 FT CARBOHYD 295 295 N-linked (GlcNAc...)
 FT CARBOHYD 301 301 N-linked (GlcNAc...)
 FT CARBOHYD 306 306 N-linked (GlcNAc...)
 FT CARBOHYD 338 338 N-linked (GlcNAc...)
 FT CARBOHYD 601 601 N-linked (GlcNAc...)
 FT CARBOHYD 621 621 N-linked (GlcNAc...)
 FT CARBOHYD 681 681 N-linked (GlcNAc...)
 FT CARBOHYD 892 892 N-linked (GlcNAc...)
 FT CARBOHYD 1060 1060 N-linked (GlcNAc...)
 FT CARBOHYD 1064 1064 N-linked (GlcNAc...)
 FT CARBOHYD 1080 1080 N-linked (GlcNAc...)
 FT CARBOHYD 1146 1146 N-linked (GlcNAc...)
 FT CARBOHYD 1378 1378 N-linked (GlcNAc...)
 FT CARBOHYD 1392 1392 N-linked (GlcNAc...)
 FT CARBOHYD 1403 1403 N-linked (GlcNAc...)
 SQ SEQUENCE 2009 AA; 228769 MW; 6808466F6368373B CRC64;

Query Match 100.0%; Score 28; DB 1; Length 2009;
 Best Local Similarity 100.0%; Pred. No. 5.5e-19;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 MYLYFYVFIIFGSPFTLNFTGVINDNF 28
 Db 1459 MYLYFYVFIIFGSPFTLNFTGVINDNF 1486

RESULT 19
 ID 090230 PRELIMINARY; PRT; 718 AA.
 AC 090230
 DT 01-DEC-2001 (TREMBlrel. 19, Created)
 DT 01-DEC-2001 (TREMBlrel. 19, Last sequence update)
 DT 01-MAR-2004 (TREMBlrel. 26, Last annotation update)
 DE Sodium channel 1 (Fragment).
 OS Sternohygus macrurus.
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Actinopterygii; Neopterygii; Teleostei; Osteichthyes; Gymnotiformes;
 OC Sternopygidae; Sternopygus.
 OX NCBI_TaxId=77841;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=21310016; PubMed=11416226;
 RA Lopreato G.F., Lu Y., Southwell A., Atkinson N.S., Hillis D.M.,
 RT Wilcox T.P., Zakon H.H.;
 RL "Evolution and divergence of sodium channel genes in vertebrates";
 DR EMBL; AF378139; AAK55437.1; -;
 DR GO; GO:0016021; C:integral to membrane; IEA.
 DR GO; GO:0005261; C:integral to membrane; IEA.
 DR GO; GO:0006812; P:cation transport; IEA.
 DR InterPro; IPR001682; Ca/Na pore.
 DR InterPro; IPR002111; Cat channel_TrpL.
 DR InterPro; IPR005821; Ion trans.
 DR InterPro; IPR005820; M-channel_nlg.
 DR InterPro; IPR010526; Na trans_assoc.
 DR Pfam; PF00520; Ion trans; 2
 DR Pfam; PF06512; Na_trans_assoc; 1.
 KW Ion transport; Ionic channel; Transmembrane; Transport.
 FT NON_TER 1 1
 FT NON_TER 718 718
 SQ SEQUENCE 718 AA; 81545 MW; 67C779B99DA3BCE CRC64;

Query Match 89.3%; Score 25; DB 2; Length 718;
 Best Local Similarity 100.0%; Pred. No. 2.2e-16;
 Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 4 YFVIFIFGSPFTLNFTGVINDNF 28
 Db 644 YFVIFIFGSPFTLNFTGVINDNF 668

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RESULT 20
O90226 PRELIMINARY; PRT; 743 AA.
ID O90226;
DT 01-DEC-2001 (TREMBlrel. 19, Created)
DT 01-DEC-2001 (TREMBlrel. 19, Last sequence update)
DT 01-MAR-2004 (TREMBlrel. 26, Last annotation update)
DE Sodium channel 5 (Fragment).
OS Sternopygus macrurus.
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Gymnotiformes;
OC Sternopygidae; Sternopygus.
OX NCBI_TaxID=77841;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=21310016; PubMed=11416226;
RA Lopreato G.F., Lu Y., Southwell A., Atkinson N.S., Hillis D.M.,
RA Wilcox T.P., Zakon H.H.;
RT "Evolution and divergence of sodium channel genes in vertebrates.";
RL Proc. Natl. Acad. Sci. U.S.A. 98:7588-7592(2001).
DR EMBL; AF378143; AAK55441.1; -.
DR GO; GO:0016021; C:Integral to membrane; IEA.
DR GO; GO:0005261; F:cation channel activity; IEA.
DR GO; GO:0006812; P:cation transport; IEA.
DR InterPro; IPR001682; Ca/Na.pore.
DR InterPro; IPR002111; Cat_channel_TrpL.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR005820; M+channel_nlg.
DR InterPro; IPR010526; Na_trans_assoc.
DR Pfam; PF00520; Ion_trans; 2.
DR Pfam; PF06512; Na_trans_assoc; 1.
DR Ion transport; Ionic channel; Transmembrane; Transport.
KW ION_TRAN
FT NON_TER
FT TER
SQ SEQUENCE 743 AA; 84781 MW; F242966554CAB0C CRC64;

Query Match 89.3%; Score 25; DB 2; Length 743;
Best Local Similarity 100.0%; Pred. No. 2,36-16;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 YFVIFIFGSPFTLNLFVGIINDNF 28
DB 669 YFVIFIFGSPFTLNLFVGIINDNF 693

RESULT 21
O91BF1 PRELIMINARY; PRT; 1880 AA.
ID O91BF1;
AC O91BF1;
DT 01-OCT-2000 (TREMBlrel. 15, Created)
DT 01-OCT-2000 (TREMBlrel. 15, Last sequence update)
DT 01-MAR-2004 (TREMBlrel. 26, Last annotation update)
DE Voltage-gated sodium channel.
OS Takifugu pardalis (Puffer fish).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Euteleostei;
OC Acanthomorphi; Acanthopterygii; Percormorpha; Tetraodontiformes;
OC Tetraodontidae; Tetraodontidae; Takifugu.
OX NCBI_TaxID=98921;
RN [1]
RP SEQUENCE FROM N.A.
RX TISSUE=Skletal muscle;
RX MEDLINE=20090650; PubMed=10623632;
RA Yocesu-Yamashita M., Nishimori K., Nitani Y., Isemura M., Sugimoto A.,
RA Yasumoto T.;
RT "Binding properties of 3H-PbTx-3 and 3H-saxitoxin to brain membranes
and to skeletal muscle membranes of puffer fish Takifugu pardalis, and the
primary structure of a voltage-gated Na+ channel alpha-subunit (fMna1)
from skeletal muscle of F. pardalis.";
RL Biochem. Biophys. Res. Commun. 267:403-412(2000).
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).

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CC -1- SIMILARITY: Belongs to the sodium channel family.
DR EMBL; AB030482; BAA30398.1; -.
DR HSSP; P04775; 1BYX.
DR GO; GO:0016021; C:Integral to membrane; IEA.
DR GO; GO:0001518; C:voltage-gated sodium channel complex; IEA.
DR GO; GO:0005261; F:cation channel activity; IEA.
DR GO; GO:0005248; F:voltage-gated sodium channel activity; IEA.
DR GO; GO:0006812; P:cation transport; IEA.
DR GO; GO:0006814; P:sodium ion transport; IEA.
DR InterPro; IPR001682; Ca/Na.pore.
DR InterPro; IPR002111; Cat_channel_TrpL.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IQ_region.
DR InterPro; IPR005820; M+channel_nlg.
DR InterPro; IPR010526; Na_channel.
DR InterPro; IPR010526; Na_trans_assoc.
DR Pfam; PF00520; Ion_trans; 4.
DR Pfam; PF00612; IQ; 1.
DR Pfam; PF06512; Na_trans_assoc; 1.
DR PRINTS; PR00170; NACHANNEL.
DR PROSITE; PS50096; IQ; 1.
KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
KW Transport; Voltage-gated channel.
SQ SEQUENCE 1880 AA; 212084 MW; 4064836C3D43E02 CRC64;

Query Match 89.3%; Score 25; DB 2; Length 1880;
Best Local Similarity 100.0%; Pred. No. 4,86-16;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 YFVIFIFGSPFTLNLFVGIINDNF 28
DB 1302 YFVIFIFGSPFTLNLFVGIINDNF 1326

RESULT 22
O9DF53 PRELIMINARY; PRT; 1949 AA.
ID O9DF53;
AC O9DF53;
DT 01-MAR-2001 (TREMBlrel. 16, Created)
DT 01-MAR-2001 (TREMBlrel. 16, Last sequence update)
DT 01-MAR-2004 (TREMBlrel. 26, Last annotation update)
DE Sodium channel protein Scn8a.
OS Brachydanio rerio (Zebrafish) (Danio rerio).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;
OC Cyprinidae; Danio.
OX NCBI_TaxID=7955;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Oregon.
RA Tsai C.-W., Tseng J.-J., Horng J.-F., Wu J.-L., Tsay H.-J.;
RL Submitted (AUG-2000) to the EMBL/GenBank/DBJ databases.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
CC -1- SIMILARITY: Belongs to the sodium channel family.
DR EMBL; AF297658; AAG18440.1; -.
DR HSSP; P04775; 1BYX.
DR ZFIN; ZDB-GENE-000828-1; scn8a.
DR GO; GO:0016021; C:Integral to membrane; IEA.
DR GO; GO:0001518; C:voltage-gated sodium channel complex; IEA.
DR GO; GO:0005261; F:cation channel activity; IEA.
DR GO; GO:0005248; F:voltage-gated sodium channel activity; IEA.
DR GO; GO:0006812; P:cation transport; IEA.
DR GO; GO:0006814; P:sodium ion transport; IEA.
DR InterPro; IPR001682; Ca/Na.pore.
DR InterPro; IPR002111; Cat_channel_TrpL.
DR InterPro; IPR000048; IQ_region.
DR InterPro; IPR005820; M+channel_nlg.
DR InterPro; IPR010526; Na_channel.
DR InterPro; IPR010526; Na_trans_assoc.
DR Pfam; PF00520; Ion_trans; 4.

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DR Pfam; PF00612; IQ; 1.
 DR Pfam; PF06512; Na_trans_assoc; 1.
 DR PRINTS; PRO0170; NACHANNEL.
 DR PRINTS; PRO1667; NACHANNEL8.
 DR SMART; SM00015; IQ; 1.
 KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
 KW Transport; Voltage-gated channel.
 SQ SEQUENCE 1949 AA; 221760 MW; 6BAC69664B0C7BC3 CRC64;

Query Match 89.3%; Score 25; DB 2; Length 1949;
 Best Local Similarity 100.0%; Pred. No. 4.9e-16;
 Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 YFVIFIFGSPFTLNLFIVGIIDNF 28
 DB 1412 YFVIFIFGSPFTLNLFIVGIIDNF 1436

RESULT 23
 ID 075RX9 PRELIMINARY; PRT; 1962 AA.
 AC 075RX9;
 DT 05-JUL-2004 (TRENBLrel. 27, Created)
 DT 05-JUL-2004 (TRENBLrel. 27, Last sequence update)
 DT 05-JUL-2004 (TRENBLrel. 27, Last annotation update)
 DE TTX-resistant sodium channel splicing variant.
 GN Name=Nav1.5;
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Ou S., Kameyama A., Kameyama M.;
 RL Submitted (JAN-2004) to the EMBL/GenBank/DBJ databases.
 CC [2]
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
 CC -1- SIMILARITY: Belongs to the sodium channel family.
 DR EMBL; AB158470; BAD12085.1;
 DR GO; GO:0005216; F:ion channel activity; IEA.
 DR InterPro; IPR001682; Ca/Na_pore.
 DR InterPro; IPR002111; Cat_channel_TrypL.
 DR InterPro; IPR010983; BF_Hand_like.
 DR InterPro; IPR005821; Ion_trans.
 DR InterPro; IPR000048; IQ_region.
 DR InterPro; IPR005820; M+channel_nlg.
 DR InterPro; IPR001696; Na_channel.
 DR InterPro; IPR008053; Na_channel5.
 DR InterPro; IPR010526; Na_trans_assoc.
 DR Pfam; PF00520; Ion_trans; 4.
 DR Pfam; PF00612; IQ; 1.
 DR Pfam; PF06512; Na_trans_assoc; 1.
 DR PRINTS; PRO0170; NACHANNEL.
 DR PRINTS; PRO1666; NACHANNEL5.
 KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
 KW Transport; Voltage-gated channel.
 SQ SEQUENCE 1962 AA; 221146 MW; 3EA3B3D897199C9A CRC64;

Query Match 89.3%; Score 25; DB 2; Length 1962;
 Best Local Similarity 100.0%; Pred. No. 5e-16;
 Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 YFVIFIFGSPFTLNLFIVGIIDNF 28
 DB 1395 YFVIFIFGSPFTLNLFIVGIIDNF 1419

RESULT 24
 BAD12085 PRELIMINARY; PRT; 1962 AA.

AC BAD12085;
 DT 03-MAR-2004 (TRENBLrel. 27, Created)
 DT 03-MAR-2004 (TRENBLrel. 27, Last sequence update)
 DT 03-MAR-2004 (TRENBLrel. 27, Last annotation update)
 DE TTX-resistant sodium channel splicing variant.
 GN NAV1.5.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Ou S., Kameyama A., Kameyama M.;
 RL Submitted (JAN-2004) to the EMBL/GenBank/DBJ databases.
 CC [2]
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
 CC -1- SIMILARITY: Belongs to the sodium channel family.
 DR EMBL; U39018; AAC42059.1; --
 DR PIR; I56555; I56555;
 DR HSP; P04775; I8Y.
 DR GO; GO:0016021; C:integral to membrane; IEA.
 DR GO; GO:001518; C:voltage-gated sodium channel complex; IEA.
 DR GO; GO:0005261; F:cation channel activity; IEA.
 DR GO; GO:0005248; F:voltage-gated sodium channel activity; IEA.
 DR GO; GO:0006814; P:cation ion transport; IEA.
 DR InterPro; IPR001682; Ca/Na_pore.
 DR InterPro; IPR002111; Cat_channel_TrypL.
 DR InterPro; IPR005821; Ion_trans.
 DR InterPro; IPR000048; IQ_region.
 DR InterPro; IPR001696; Na_channel_nlg.
 DR InterPro; IPR005820; M+channel.
 DR InterPro; IPR008054; Na_channel8.

Query Match 89.3%; Score 25; DB 2; Length 1962;
 Best Local Similarity 100.0%; Pred. No. 5e-16;
 Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 YFVIFIFGSPFTLNLFIVGIIDNF 28
 DB 1395 YFVIFIFGSPFTLNLFIVGIIDNF 1419

RESULT 25
 ID 063541 PRELIMINARY; PRT; 1976 AA.
 AC 063541;
 DT 01-NOV-1996 (TRENBLrel. 01, Created)
 DT 01-NOV-1996 (TRENBLrel. 01, Last sequence update)
 DT 01-MAR-2004 (TRENBLrel. 26, Last annotation update)
 DE Sodium channel protein 6.
 GN Name=SCP6;
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 OX NCBI_TaxID=10116;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=Sprague-Dawley; TISSUE=Brain;
 RX MEDLINE=95271284; PubMed=7751906;
 RA Schaller K.L., Krzemien D.M., Yarowsky P.J., Krueger B.K.,
 RA Caldwell J.H.;
 RT "A novel, abundant sodium channel expressed in neurons and glia."
 RL J. Neurosci. 15:3231-3242(1995).
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
 CC -1- SIMILARITY: Belongs to the sodium channel family.
 DR EMBL; U39018; AAC42059.1; --
 DR PIR; I56555; I56555;
 DR HSP; P04775; I8Y.
 DR GO; GO:0016021; C:integral to membrane; IEA.
 DR GO; GO:001518; C:voltage-gated sodium channel complex; IEA.
 DR GO; GO:0005261; F:cation channel activity; IEA.
 DR GO; GO:0005248; F:voltage-gated sodium channel activity; IEA.
 DR GO; GO:0006814; P:cation ion transport; IEA.
 DR InterPro; IPR001682; Ca/Na_pore.
 DR InterPro; IPR002111; Cat_channel_TrypL.
 DR InterPro; IPR005821; Ion_trans.
 DR InterPro; IPR000048; IQ_region.
 DR InterPro; IPR001696; Na_channel_nlg.
 DR InterPro; IPR005820; M+channel.
 DR InterPro; IPR008054; Na_channel8.

DR InterPro; IPR010526; Na_trans_assoc.
 DR Pfam; PF00520; Ion_trans; 4.
 DR Pfam; PF00612; IQ; 1.
 DR PRINTS; PRO0170; Na_trans_assoc; 1.
 DR PRINTS; PRO0170; NACHANNEL.
 DR SMART; SM00015; IQ; 1.
 DR PROSITE; PS00096; IQ; 1.
 DR PROSITE; PS00878; ODR_DC_2_1; UNKNOWN_1.
 KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
 KW Transport; Voltage-gated channel.
 SQ SEQUENCE 1976 AA; 225227 MW; B6949327A47FA88A CRC64;

Query Match 89.3%; Score 25; DB 2; Length 1976;
 Best Local Similarity 100.0%; Pred.No. 5e-16;
 Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 YFVIFIIFGSFTLNLFIGVIIDNP 28
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 DB 1439 YFVIFIIFGSFTLNLFIGVIIDNP 1463

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OW protein - protein search, using sw model

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- 6: /cgn2_6/ptodata/1/1aa/backfills1.pep:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	28	100.0	1836	4 US-10-162-012-24	Sequence 24, Appl
2	28	100.0	2005	3 US-08-836-325-7	Sequence 7, Appl
3	28	100.0	2005	4 US-09-457-571-7	Sequence 7, Appl
4	25	89.3	813	3 US-08-836-325-8	Sequence 8, Appl
5	25	89.3	813	4 US-09-457-571-8	Sequence 8, Appl
6	25	89.3	1011	3 US-08-836-325-2	Sequence 2, Appl
7	25	89.3	1011	4 US-09-457-571-2	Sequence 2, Appl
8	25	89.3	1976	3 US-09-024-020B-9	Sequence 9, Appl
9	25	89.3	1976	3 US-09-425-043-9	Sequence 9, Appl
10	25	89.3	1978	3 US-09-024-020B-3	Sequence 3, Appl
11	25	89.3	1978	3 US-09-425-043-3	Sequence 3, Appl
12	25	89.3	1984	3 US-08-836-325-10	Sequence 10, Appl
13	25	89.3	1984	4 US-09-457-571-10	Sequence 10, Appl
14	25	89.3	1988	3 US-09-024-020B-4	Sequence 4, Appl
15	25	89.3	1988	3 US-09-425-043-4	Sequence 4, Appl
16	25	89.3	1989	3 US-08-836-325-12	Sequence 12, Appl
17	25	89.3	1989	4 US-09-457-571-12	Sequence 12, Appl
18	25	89.3	2016	3 US-09-634-920-4	Sequence 4, Appl
19	25	89.3	2016	4 US-09-514-907A-2	Sequence 2, Appl
20	25	89.3	2016	4 US-09-896-994-2	Sequence 2, Appl
21	25	89.3	2016	4 US-09-840-125-4	Sequence 4, Appl
22	21	75.0	413	2 US-08-808-793-25	Sequence 25, Appl
23	21	75.0	452	3 US-07-998-289B-6	Sequence 6, Appl
24	21	75.0	1233	4 US-09-354-147C-7	Sequence 7, Appl
25	21	75.0	1243	4 US-09-354-147C-8	Sequence 8, Appl
26	21	75.0	1765	4 US-09-354-147C-2	Sequence 2, Appl
27	21	75.0	1765	4 US-09-354-147C-3	Sequence 3, Appl

28	21	75.0	1765	4 US-09-354-147C-5	Sequence 5, Appl
29	21	75.0	1791	4 US-09-354-147C-42	Sequence 42, Appl
30	21	75.0	1820	3 US-07-998-289B-8	Sequence 8, Appl
31	21	75.0	1835	3 US-08-836-325-15	Sequence 15, Appl
32	21	75.0	1835	4 US-09-457-571-15	Sequence 15, Appl
33	21	75.0	1835	4 US-08-836-325-16	Sequence 16, Appl
34	21	75.0	1969	4 US-09-457-571-16	Sequence 16, Appl
35	21	75.0	1977	4 US-09-976-594-757	Sequence 757, App
36	21	75.0	1977	4 US-09-919-039-367	Sequence 367, App
37	21	75.0	1989	3 US-08-836-325-11	Sequence 11, Appl
38	21	75.0	1989	4 US-09-457-571-11	Sequence 11, Appl
39	21	75.0	2100	2 US-08-808-793-23	Sequence 23, Appl
40	21	75.0	2100	3 US-08-772-512A-19	Sequence 19, Appl
41	21	75.0	2104	2 US-08-808-793-4	Sequence 4, Appl
42	21	75.0	2104	3 US-08-772-512A-4	Sequence 4, Appl
43	21	75.0	2105	2 US-08-808-793-3	Sequence 3, Appl
44	21	75.0	2105	3 US-08-772-512A-3	Sequence 3, Appl
45	12	42.9	1956	3 US-08-843-417-10	Sequence 10, Appl
46	12	42.9	1956	4 US-09-527-013-10	Sequence 10, Appl
47	7	25.0	8	3 US-08-836-325-6	Sequence 6, Appl
48	7	25.0	8	4 US-09-457-571-6	Sequence 6, Appl
49	7	25.0	163	4 US-09-134-000C-5625	Sequence 5625, Ap
50	7	25.0	307	3 US-08-605-284B-19	Sequence 19, Appl
51	7	25.0	309	3 US-08-605-284B-11	Sequence 11, Appl
52	7	25.0	310	3 US-08-605-284B-10	Sequence 10, Appl
53	7	25.0	310	3 US-08-605-284B-12	Sequence 12, Appl
54	7	25.0	310	3 US-08-605-284B-13	Sequence 13, Appl
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56	7	25.0	310	3 US-08-605-284B-15	Sequence 15, Appl
57	7	25.0	311	3 US-08-605-284B-18	Sequence 18, Appl
58	7	25.0	311	3 US-08-605-284B-23	Sequence 23, Appl
59	7	25.0	312	3 US-08-605-284B-20	Sequence 20, Appl
60	7	25.0	434	3 US-08-489-039A-13633	Sequence 13633, A
61	7	25.0	1956	3 US-08-843-417-2	Sequence 2, Appl
62	7	25.0	1956	4 US-09-527-013-2	Sequence 2, Appl
63	7	25.0	1957	4 US-08-669-656A-2	Sequence 2, Appl
64	7	25.0	1957	4 US-08-669-656A-8	Sequence 8, Appl
65	7	25.0	2132	4 US-08-669-656A-6	Sequence 6, Appl
66	7	25.0	222	1 US-08-103-445-12	Sequence 12, Appl
67	6	21.4	22	1 US-08-461-660B-12	Sequence 12, Appl
68	6	21.4	22	1 US-09-270-767-61121	Sequence 61121, A
69	6	21.4	55	4 US-09-328-352-4945	Sequence 4945, Ap
70	6	21.4	63	4 US-09-107-532A-4445	Sequence 4445, Ap
71	6	21.4	70	4 US-09-134-000C-4125	Sequence 4125, Ap
72	6	21.4	130	3 US-09-134-000C-5023	Sequence 5023, Ap
73	6	21.4	143	4 US-09-270-767-38745	Sequence 38745, A
74	6	21.4	143	4 US-09-270-767-53962	Sequence 53962, A
75	6	21.4	180	4 US-09-270-767-38105	Sequence 38105, A
76	6	21.4	180	4 US-09-270-767-53322	Sequence 53322, A
77	6	21.4	237	4 US-09-538-032-332	Sequence 332, App
78	6	21.4	261	4 US-09-328-352-4430	Sequence 4430, App
79	6	21.4	262	4 US-09-489-039A-12430	Sequence 12430, A
80	6	21.4	262	4 US-09-543-681A-6127	Sequence 6127, Ap
81	6	21.4	274	4 US-09-134-000C-6450	Sequence 6450, Ap
82	6	21.4	298	3 US-09-134-001C-3631	Sequence 3631, Ap
83	6	21.4	316	2 US-08-846-762-9	Sequence 9, Appl
84	6	21.4	318	4 US-09-710-279-3118	Sequence 3118, Ap
85	6	21.4	322	3 US-09-134-001C-4064	Sequence 4064, Ap
86	6	21.4	326	4 US-09-107-532A-3665	Sequence 3665, Ap
87	6	21.4	404	4 US-09-543-681A-6702	Sequence 6702, Ap
88	6	21.4	421	3 US-09-134-001C-4890	Sequence 4890, Ap
89	6	21.4	426	4 US-09-491-577-46	Sequence 46, Appl
90	6	21.4	430	4 US-09-489-039A-10587	Sequence 10587, A
91	6	21.4	435	4 US-09-275-252A-5	Sequence 5, Appl
92	6	21.4	495	2 US-08-527-152-2	Sequence 2, Appl
93	6	21.4	528	4 US-09-252-991A-20769	Sequence 20769, A
94	6	21.4	672	4 US-09-556-916-26	Sequence 26, Appl
95	6	21.4	672	4 US-09-556-916-28	Sequence 28, Appl

ALIGNMENTS

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RESULT 1
US-10-162-012-24
; Sequence 24, Application US/10162012
; Patent No. 6682597
; GENERAL INFORMATION:
; APPLICANT: Curtis, Rory A.J.
; APPLICANT: Silos-Santiago, Immaculada
; APPLICANT: Gu, Wei
; TITLE OF INVENTION: NOVEL HUMAN ION CHANNEL AND TRANSPORTER FAMILY MEMBERS
; FILE REFERENCE: 10448-190001
; CURRENT FILING DATE: US/10/162,012
; PRIOR FILING DATE: 2002-06-04
; PRIOR APPLICATION NUMBER: US 60/209,845
; PRIOR FILING DATE: 2000-06-06
; PRIOR APPLICATION NUMBER: US 09/875,321
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: PCT/US01/18340
; PRIOR FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: US 60/209,257
; PRIOR FILING DATE: 2000-06-05
; PRIOR APPLICATION NUMBER: US 09/875,423
; PRIOR FILING DATE: 2001-06-05
; PRIOR APPLICATION NUMBER: PCT/US01/18398
; PRIOR FILING DATE: 2001-06-05
; PRIOR APPLICATION NUMBER: US 60/209,238
; PRIOR FILING DATE: 2000-06-05
; PRIOR APPLICATION NUMBER: US 09/875,363
; PRIOR FILING DATE: 2001-06-05
; PRIOR APPLICATION NUMBER: PCT/US01/18247
; PRIOR FILING DATE: 2001-06-05
; PRIOR APPLICATION NUMBER: US 60/227,068
; PRIOR FILING DATE: 2000-08-22
; PRIOR APPLICATION NUMBER: US 09/928,530
; PRIOR FILING DATE: 2001-08-13
; PRIOR APPLICATION NUMBER: PCT/US01/25475
; PRIOR FILING DATE: 2001-08-15
; PRIOR APPLICATION NUMBER: US 60/226,770
; PRIOR FILING DATE: 2000-08-21
; PRIOR APPLICATION NUMBER: US 09/934,421
; PRIOR FILING DATE: 2001-08-21
; PRIOR APPLICATION NUMBER: PCT/US01/26096
; PRIOR FILING DATE: 2001-08-21
; PRIOR APPLICATION NUMBER: US 60/279,281
; PRIOR FILING DATE: 2001-03-28
; PRIOR APPLICATION NUMBER: US 10/109,029
; PRIOR FILING DATE: 2002-03-28
; PRIOR APPLICATION NUMBER: PCT/US02/09728
; PRIOR FILING DATE: 2002-03-28
; PRIOR APPLICATION NUMBER: US 60/290,288
; PRIOR FILING DATE: 2001-05-11
; PRIOR APPLICATION NUMBER: US (not assigned)
; PRIOR FILING DATE: 2002-05-13
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 24
; LENGTH: 1836
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-162-012-24

Query Match      100.0%; Score 28; DB 4; Length 1836;
Best Local Similarity 100.0%; Pred. No. 1.6e-20;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY      1 MYLVFVIFIIIGSFPTLNLFIGVIIDNF 28
DB      1271 MYLVFVIFIIIGSFPTLNLFIGVIIDNF 1298

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RESULT 2
US-08-836-325-7
; Sequence 7, Application US/08836325
; Patent No. 6110672
; GENERAL INFORMATION:
; APPLICANT: Mandel, Gail
; APPLICANT: Halegoua, Simon
; APPLICANT: Borden, Laurence A.
; TITLE OF INVENTION: Peripheral Nervous System Specific
; TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
; TITLE OF INVENTION: X-Ray Diffraction, Computer Molecular Modeling, Rational
; TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
; TITLE OF INVENTION: Thereof
; NUMBER OF SEQUENCES: 19
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
; STREET: 1100 New York Ave., N.W., Suite 600
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20005-3934
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: IBM PC compatible
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/836,325
; FILING DATE: 2-MAY-1997
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/14251
; FILING DATE: 02-NOV-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/482,401
; FILING DATE: 07-JUN-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/334,029
; FILING DATE: 02-NOV-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Ludwig, Steven R.
; REGISTRATION NUMBER: 36,203
; REFERENCE/DOCKET NUMBER: 0917.0240002
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-371-2600
; TELEFAX: 202-371-2540
; INFORMATION FOR SEQ ID NO: 7:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2005 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
US-08-836-325-7

Query Match      100.0%; Score 28; DB 3; Length 2005;
Best Local Similarity 100.0%; Pred. No. 1.7e-20;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY      1 MYLVFVIFIIIGSFPTLNLFIGVIIDNF 28
DB      1449 MYLVFVIFIIIGSFPTLNLFIGVIIDNF 1476

RESULT 3
US-09-457-571-7
; Sequence 7, Application US/09457571
; Patent No. 6703486
; GENERAL INFORMATION:
; APPLICANT: Mandel, Gail
; APPLICANT: Halegoua, Simon
; TITLE OF INVENTION: Peripheral Nervous System Specific

```

TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
TITLE OF INVENTION: Thereof
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/457,571
FILING DATE: 09-DEC-1999
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/836,325
FILING DATE: 02-MAY-1997
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917,0240003
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 2005 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-457-571-7

Query Match 100.0%; Score 28; DB 4; Length 2005;
Best Local Similarity 100.0%; Pred. No. 1.7e-20;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 MYLVFVFIIFGSPFTLNLFGVIIDNF 28
Db 1449 MYLVFVFIIFGSPFTLNLFGVIIDNF 1476

RESULT 4
US-08-836-325-8
Sequence 8, Application US/08836325
Patent No. 6110672
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Halegoua, Simon
APPLICANT: Borden, Laurence A.
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:

ADDRESSEE: STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/836,325
FILING DATE: 2-MAY-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917,0240002
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 8:
SEQUENCE CHARACTERISTICS:
LENGTH: 813 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-08-836-325-8

Query Match 89.3%; Score 25; DB 3; Length 813;
Best Local Similarity 100.0%; Pred. No. 8.9e-18;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 4 YFVFIIFGSPFTLNLFGVIIDNF 28
Db 347 YFVFIIFGSPFTLNLFGVIIDNF 371

RESULT 5
US-09-457-571-8
Sequence 8, Application US/09457571
Patent No. 6703486
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Halegoua, Simon
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS

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SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/457,571
FILING DATE: 09-DEC-1999
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/836,325
FILING DATE: 02-MAY-1997
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917.0240003
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 8:
SEQUENCE CHARACTERISTICS:
LENGTH: 813 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-457-571-8

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Query Match      89.3%; Score 25; DB 4; Length 813;
Best Local Similarity 100.0%; Pred. No. 8.9e-18;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 4 YFVFIIFGSPFTLNLFIVGIIDNF 28
Db 347 YFVFIIFGSPFTLNLFIVGIIDNF 371

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RESULT 6
US-08-836-325-2
Sequence 2, Application US/08836325
Patent No. 6110672
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Halegoua, Simon
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational,
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSER: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N.W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/836,325
FILING DATE: 2-MAY-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:

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APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917.0240002
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 1011 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-836-325-2

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Query Match      89.3%; Score 25; DB 3; Length 1011;
Best Local Similarity 100.0%; Pred. No. 1.1e-17;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 4 YFVFIIFGSPFTLNLFIVGIIDNF 28
Db 481 YFVFIIFGSPFTLNLFIVGIIDNF 505

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RESULT 7
US-09-457-571-2
Sequence 2, Application US/09457571
Patent No. 6703486
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Halegoua, Simon
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational,
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSER: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N.W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/457,571
FILING DATE: 09-DEC-1999
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/836,325
FILING DATE: 02-MAY-1997
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994

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ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917, 0240003
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 1011 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-09-457-571-2

Query Match 89.3%; Score 25; DB 4; Length 1011;
Best Local Similarity 100.0%; Pred. No. 1.1e-17;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 4 YFVFIIFGSPFTLNLFIVGIINDF 28
DB 481 YFVFIIFGSPFTLNLFIVGIINDF 505

RESULT 8
US-09-024-0208-9
Sequence 9, Application US/090240208
Patent No. 6030810
GENERAL INFORMATION:
APPLICANT: DELGADO, STEPHEN G.
APPLICANT: DIETRICH, PAUL S.
APPLICANT: FISH, LINDA M.
APPLICANT: HERMAN, RONALD C.
APPLICANT: SANGAMESWARAN, LAKSHMI
TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE
NUMBER OF SEQUENCES: 43
CURRENT APPLICATION DATA:
CORRESPONDENCE ADDRESS:
ADDRESSEE: JANET PAULINE CLARK
STREET: 3401 HILLVIEW AVENUE, MS A2-250
CITY: PALO ALTO
STATE: CA
COUNTRY: U.S.A.
ZIP: 94304-1397
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/024,0208
FILING DATE: 16-FEB-1998
CLASSIFICATION: 536
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/039,447
FILING DATE: 26-FEB-1997
ATTORNEY/AGENT INFORMATION:
NAME: CLARK, JANET P.
REGISTRATION NUMBER: 34,799
REFERENCE/DOCKET NUMBER: R00208-REG
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 852-3097
TELEFAX: (650) 855-5322
INFORMATION FOR SEQ ID NO: 9:
SEQUENCE CHARACTERISTICS:
LENGTH: 1976 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-024-0208-9

Query Match 89.3%; Score 25; DB 3; Length 1976;

Best Local Similarity 100.0%; Pred. No. 1.9e-17;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 YFVFIIFGSPFTLNLFIVGIINDF 28
DB 1439 YFVFIIFGSPFTLNLFIVGIINDF 1463

RESULT 9
US-09-425-043-9
Sequence 9, Application US/09425043
Patent No. 6335172
GENERAL INFORMATION:
APPLICANT: DELGADO, STEPHEN G.
APPLICANT: DIETRICH, PAUL S.
APPLICANT: FISH, LINDA M.
APPLICANT: HERMAN, RONALD C.
APPLICANT: SANGAMESWARAN, LAKSHMI
TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE
NUMBER OF SEQUENCES: 43
CURRENT APPLICATION DATA:
CORRESPONDENCE ADDRESS:
ADDRESSEE: JANET PAULINE CLARK
STREET: 3401 HILLVIEW AVENUE, MS A2-250
CITY: PALO ALTO
STATE: CA
COUNTRY: U.S.A.
ZIP: 94304-1397
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/425,043
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 09/024,020
FILING DATE: 16-FEB-1998
APPLICATION NUMBER: US 60/039,447
FILING DATE: 26-FEB-1997
ATTORNEY/AGENT INFORMATION:
NAME: CLARK, JANET P.
REGISTRATION NUMBER: 34,799
REFERENCE/DOCKET NUMBER: R00208-REG
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 852-3097
TELEFAX: (650) 855-5322
INFORMATION FOR SEQ ID NO: 9:
SEQUENCE CHARACTERISTICS:
LENGTH: 1976 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-425-043-9

Query Match 89.3%; Score 25; DB 3; Length 1976;
Best Local Similarity 100.0%; Pred. No. 1.9e-17;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 YFVFIIFGSPFTLNLFIVGIINDF 28
DB 1439 YFVFIIFGSPFTLNLFIVGIINDF 1463

RESULT 10
US-09-024-0208-3
Sequence 3, Application US/090240208
Patent No. 6030810
GENERAL INFORMATION:
APPLICANT: DELGADO, STEPHEN G.

APPLICANT: DIETRICH, PAUL S.
APPLICANT: FISH, LINDA M.
APPLICANT: HERMAN, RONALD C.
APPLICANT: SANGAMESWARAN, LAKSHMI
TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE
TITLE OF INVENTION: SODIUM CHANNEL I-SUBUNIT AND A SPLICE VARIANT THEREOF
NUMBER OF SEQUENCES: 43
CORRESPONDENCE ADDRESS:
ADDRESSEE: JANET PAULINE CLARK
STREET: 3401 HILLVIEW AVENUE, MS A2-250
CITY: PALO ALTO
STATE: CA
COUNTRY: U.S.A.
ZIP: 94304-1397
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/024,020B
FILING DATE: 16-FEB-1998
CLASSIFICATION: 536
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/039,447
FILING DATE: 26-FEB-1997
ATTORNEY/AGENT INFORMATION:
NAME: CLARK, JANET P.
REGISTRATION NUMBER: 34,799
REFERENCE/DOCKET NUMBER: R0020B-REG
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 852-3097
TELEFAX: (650) 855-5322
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 1978 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-024-020B-3

Query Match 89.3%; Score 25; DB 3; Length 1978;
Best Local Similarity 100.0%; Pred. No. 1.9e-17;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 4 YFVFIIFGSPFTLNLFVGIIDNF 28
Db 1441 YFVFIIFGSPFTLNLFVGIIDNF 1465

RESULT 11
US-09-425-043-3
Sequence 3, Application US/09425043
Patent No. 6335172
GENERAL INFORMATION:
APPLICANT: DELGADO, STEPHEN G.
APPLICANT: DIETRICH, PAUL S.
APPLICANT: FISH, LINDA M.
APPLICANT: HERMAN, RONALD C.
APPLICANT: SANGAMESWARAN, LAKSHMI
TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE
TITLE OF INVENTION: SODIUM CHANNEL I-SUBUNIT AND A SPLICE VARIANT THEREOF
NUMBER OF SEQUENCES: 43
CORRESPONDENCE ADDRESS:
ADDRESSEE: JANET PAULINE CLARK
STREET: 3401 HILLVIEW AVENUE, MS A2-250
CITY: PALO ALTO
STATE: CA
COUNTRY: U.S.A.
ZIP: 94304-1397
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/425,043
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 09/024,020
FILING DATE: 16-FEB-1998
APPLICATION NUMBER: US 60/039,447
FILING DATE: 26-FEB-1997
ATTORNEY/AGENT INFORMATION:
NAME: CLARK, JANET P.
REGISTRATION NUMBER: 34,799
REFERENCE/DOCKET NUMBER: R0020B-REG
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 852-3097
TELEFAX: (650) 855-5322
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 1978 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-425-043-3

Query Match 89.3%; Score 25; DB 3; Length 1978;
Best Local Similarity 100.0%; Pred. No. 1.9e-17;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 4 YFVFIIFGSPFTLNLFVGIIDNF 28
Db 1441 YFVFIIFGSPFTLNLFVGIIDNF 1465

RESULT 12
US-08-836-325-10
Sequence 10, Application US/08836325
Patent No. 6110672
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Halegoua, Simon
APPLICANT: Borden, Laurence A.
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Thereof, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C.
STREET: 1100 New York Ave., N.W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/836,325
FILING DATE: 2-MAY-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995

PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917.0240002
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 10:
SEQUENCE CHARACTERISTICS:
LENGTH: 1984 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-836-325-10

Query Match 89.3%; Score 25; DB 3; Length 1984;
Best Local Similarity 100.0%; Pred. No. 1.9e-17;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 YFVIFIFGSPFTLNLFIGVIIDNF 28
Db 1434 YFVIFIFGSPFTLNLFIGVIIDNF 1458

RESULT 13
US-09-457-571-10
Sequence 10, Application US/09457571
Patent No. 6703486
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/457,571
FILING DATE: 09-DEC-1999
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/836,325
FILING DATE: 02-MAY-1997
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917.0240003
TELECOMMUNICATION INFORMATION:

TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 10:
SEQUENCE CHARACTERISTICS:
LENGTH: 1984 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-09-457-571-10

Query Match 89.3%; Score 25; DB 4; Length 1984;
Best Local Similarity 100.0%; Pred. No. 1.9e-17;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 YFVIFIFGSPFTLNLFIGVIIDNF 28
Db 1434 YFVIFIFGSPFTLNLFIGVIIDNF 1458

RESULT 14
US-09-024-020B-4
Sequence 4, Application US/09024020B
Patent No. 6030810
GENERAL INFORMATION:
APPLICANT: DELGADO, STEPHEN G.
APPLICANT: DIETRICH, PAUL S.
APPLICANT: FISH, LINDA M.
APPLICANT: HERMAN, RONALD C.
TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE
TITLE OF INVENTION: SODIUM CHANNEL 1-SUBUNIT AND A SPLICE VARIANT THEREOF
NUMBER OF SEQUENCES: 43
CORRESPONDENCE ADDRESS:
ADDRESSEE: JANET PAULINE CLARK
STREET: 3401 HILLVIEW AVENUE, MS A2-250
CITY: PALO ALTO
STATE: CA
COUNTRY: U.S.A.
ZIP: 94304-1397
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/024,020B
FILING DATE: 16-FEB-1998
CLASSIFICATION: 536
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/039,447
FILING DATE: 26-FEB-1997
ATTORNEY/AGENT INFORMATION:
NAME: CLARK, JANET P.
REGISTRATION NUMBER: 34,799
REFERENCE/DOCKET NUMBER: R0020B-REG
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 852-3097
TELEFAX: (650) 855-5322
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 1988 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-024-020B-4

Query Match 89.3%; Score 25; DB 3; Length 1988;
Best Local Similarity 100.0%; Pred. No. 1.9e-17;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 YFVIFIFGSPFTLNLFIGVIIDNF 28

Db 1451 YFVIFIFGSFPTLNLFIGVIIDNF 1475

RESULT 15

US-09-425-043-4
; Sequence 4, Application US/09425043
; Patent No. 6335172
; GENERAL INFORMATION:
; APPLICANT: DELGADO, STEPHEN G.
; APPLICANT: DIETRICH, PAUL S.
; APPLICANT: RISH, LINDA M.
; APPLICANT: HERMAN, RONALD C.
; APPLICANT: SANGAMESWARAN, LAKSHMI
; TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE
; TITLE OF INVENTION: SODIUM CHANNEL I-SUBUNIT AND A SPLICE VARIANT THEREOF
; NUMBER OF SEQUENCES: 43
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: JANET PAULINE CLARK
; STREET: 3401 HILLVIEW AVENUE, MS A2-250
; CITY: PALO ALTO
; STATE: CA
; COUNTRY: U.S.A.
; ZIP: 94304-1397
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: IBM PC compatible
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/425,043
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 09/024,020
; FILING DATE: 16-FEB-1998
; APPLICATION NUMBER: US 60/039,447
; FILING DATE: 26-FEB-1997
; ATTORNEY/AGENT INFORMATION:
; NAME: CLARK, JANET P.
; REGISTRATION NUMBER: 34,799
; REFERENCE/DOCKET NUMBER: R0020B-REG
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (650) 852-3097
; TELEFAX: (650) 855-5322
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1988 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; US-09-425-043-4

Query Match 89.3%; Score 25; DB 3; Length 1988;
Best Local Similarity 100.0%; Pred. No. 1.9e-17;

Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 YFVIFIFGSFPTLNLFIGVIIDNF 28
Db 1451 YFVIFIFGSFPTLNLFIGVIIDNF 1475

RESULT 16

US-08-836-325-12
; Sequence 12, Application US/08836325
; Patent No. 6110672
; GENERAL INFORMATION:
; APPLICANT: Mandel, Gail
; APPLICANT: Halegoua, Simon
; APPLICANT: Borden, Laurence A.
; TITLE OF INVENTION: Peripheral Nervous System Specific
; TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
; TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational

; TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
; TITLE OF INVENTION: Thereof
; NUMBER OF SEQUENCES: 19
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
; STREET: 1100 New York Ave., N. W., Suite 600
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20005-3934

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30

CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/836,325
FILING DATE: 2-MAY-1997

CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251

FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401

FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029

FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.

REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917.0240002
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600

TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 12:
SEQUENCE CHARACTERISTICS:
LENGTH: 1989 amino acids

TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: protein

MOLECULE TYPE: protein
US-08-836-325-12

Query Match 89.3%; Score 25; DB 3; Length 1989;
Best Local Similarity 100.0%; Pred. No. 1.9e-17;

Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 YFVIFIFGSFPTLNLFIGVIIDNF 28
Db 1436 YFVIFIFGSFPTLNLFIGVIIDNF 1460

RESULT 17
US-09-457-571-12
; Sequence 12, Application US/09457571
; Patent No. 6703486
; GENERAL INFORMATION:
; APPLICANT: Mandel, Gail
; APPLICANT: Halegoua, Simon
; TITLE OF INVENTION: Peripheral Nervous System Specific
; TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
; TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
; TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using

NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
OPERATING SYSTEM: IBM PC compatible
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/457,571
FILING DATE: 09-DEC-1999
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/836,325
FILING DATE: 02-MAY-1997
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917,024003
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 12:
SEQUENCE CHARACTERISTICS:
LENGTH: 1989 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: not relevant
MOLECULE TYPE: protein
US-09-457-571-12

Query Match 89.3%; Score 25; DB 4; Length 1989;
Best Local Similarity 100.0%; Pred. No. 1.9e-17;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 4 YVFIFIFGSFTLNLFIGVIIDNF 28
Db 1436 YVFIFIFGSFTLNLFIGVIIDNF 1460

RESULT 18
US-09-634-920-4
Sequence 4, Application US/09634920
Patent No. 6342357
GENERAL INFORMATION:
APPLICANT: Splawski, Igor
APPLICANT: Keating, Mark T.
TITLE OF INVENTION: ALTERATIONS IN THE LONG QT SYNDROME GENES KVLQT1 AND
TITLE OF INVENTION: SCNSA AND METHODS FOR DETECTING SAME
FILE REFERENCE: 2323-155
CURRENT APPLICATION NUMBER: US/09/634,920
PRIOR FILING DATE: 2000-08-09
PRIOR APPLICATION NUMBER: 60/190,057
PRIOR FILING DATE: 2000-03-17
PRIOR APPLICATION NUMBER: 60/147,488
PRIOR FILING DATE: 1999-08-09
NUMBER OF SEQ ID NOS: 4
SOFTWARE: Patentin Ver. 2.0
SEQ ID NO 4
LENGTH: 2016
TYPE: PRT
ORGANISM: Homo sapiens
US-09-634-920-4

Query Match 89.3%; Score 25; DB 3; Length 2016;
Best Local Similarity 100.0%; Pred. No. 1.9e-17;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 4 YVFIFIFGSFTLNLFIGVIIDNF 28
Db 1449 YVFIFIFGSFTLNLFIGVIIDNF 1473

RESULT 19
US-09-514-907A-2
Sequence 2, Application US/09514907A
Patent No. 6567705
GENERAL INFORMATION:
APPLICANT: Kenneth B. Stokes
Jos, e Morissette
TITLE OF INVENTION: SYSTEMS FOR ENHANCING CARDIAC SIGNAL
SENSING BY CARDIAC PACEMAKERS THROUGH
GENETIC TREATMENT
NUMBER OF SEQUENCES: 12
CORRESPONDENCE ADDRESS:
ADDRESSEE: Woodcock Washburn Kurtz Mackiewicz & No. 6567705xris LLP
STREET: One Liberty Place - 46th floor
CITY: Philadelphia
STATE: PA
COUNTRY: U.S.A.
ZIP: 19103
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
OPERATING SYSTEM: IBM PC compatible
SOFTWARE: Wordperfect 6.1
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/514,907A
FILING DATE: 08-Feb-2000
CLASSIFICATION: <Unknown>
ATTORNEY/AGENT INFORMATION:
NAME: Paul K. Legaard
REGISTRATION NUMBER: 38,534
REFERENCE/DOCKET NUMBER: MEDT-0029/P-3586
TELECOMMUNICATION INFORMATION:
TELEPHONE: (215) 568-3100
TELEFAX: (215) 568-3439
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 2016 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: unknown
SEQUENCE DESCRIPTION: SEQ ID NO: 2:
US-09-514-907A-2

Query Match 89.3%; Score 25; DB 4; Length 2016;
Best Local Similarity 100.0%; Pred. No. 1.9e-17;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 4 YVFIFIFGSFTLNLFIGVIIDNF 28
Db 1449 YVFIFIFGSFTLNLFIGVIIDNF 1473

RESULT 20
US-09-696-994-2
Sequence 2, Application US/09896994
Patent No. 6665563
GENERAL INFORMATION:
APPLICANT: Ken Stokes
Jos e Morissette
TITLE OF INVENTION: SYSTEMS AND METHODS FOR ENHANCING CARDIAC
SIGNAL SENSING BY CARDIAC PACEMAKERS THROUGH GENETIC TREATM
CORRESPONDENCE ADDRESS:
ADDRESSEE: Woodcock Washburn Kurtz Mackiewicz and No. 6665563xris
STREET: One Liberty Place - 46th floor
CITY: Philadelphia
STATE: PA

```

; COUNTRY: U.S.A.
; ZIP: 19103
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: WordPerfect 6.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/896,994
; FILING DATE: 02-Jul-2001
; CLASSIFICATION: <unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 09/514,907
; FILING DATE: <unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Paul K. Legard
; REGISTRATION NUMBER: 38,534
; REFERENCE/DOCKET NUMBER: MEDT-0029/P-3586
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (215) 568-3100
; TELEFAX: (215) 568-3439
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2016 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: unknown
; US-09-896-994-2
; SEQUENCE DESCRIPTION: SEQ ID NO: 2:

```

Query Match 89.3%; Score 25; DB 4; Length 2016;
 Best Local Similarity 100.0%; Pred. No. 1.9e-17;
 Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1449 YFVIFIFGSPFTLNLFIVGIIDNF 1473

```

RESULT 21
US-09-840-125-4
; Sequence 4, Application US/09840125
; Patent No. 6787309
; GENERAL INFORMATION:
; APPLICANT: Splawski, Igor
; TITLE OF INVENTION: Keating, Mark T.
; TITLE OF INVENTION: ALTERATIONS IN THE LONG QT SYNDROME GENES KVLQT1 AND
; FILE REFERENCE: 2323-155
; CURRENT APPLICATION NUMBER: US/09/840,125
; CURRENT FILING DATE: 2001-04-24
; PRIOR APPLICATION NUMBER: 09/634,920
; PRIOR FILING DATE: 2000-08-09
; PRIOR APPLICATION NUMBER: 60/147,488
; PRIOR FILING DATE: 1999-08-09
; NUMBER OF SEQ ID NOS: 4
; SOFTWARE: Patent In Ver. 2.0
; SEQ ID NO 4
; LENGTH: 2016
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-840-125-4

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Query Match 89.3%; Score 25; DB 4; Length 2016;
 Best Local Similarity 100.0%; Pred. No. 1.9e-17;
 Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1449 YFVIFIFGSPFTLNLFIVGIIDNF 1473

RESULT 22

```

US-08-808-793-25
; Sequence 25, Application US/08808793
; Patent No. 5858713
; GENERAL INFORMATION:
; APPLICANT: Soderlund, David M.
; APPLICANT: Ingles, Patricia J.
; TITLE OF INVENTION: CALCIUM PERMEABLE INSECT SODIUM CHANNELS
; TITLE OF INVENTION: AND USE THEREOF
; NUMBER OF SEQUENCES: 32
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Nixon, Hargrave, Devans & Doyle LLP
; STREET: Clinton Square, P.O. Box 1051
; CITY: Rochester
; STATE: New York
; COUNTRY: USA
; ZIP: 14603
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/808,793
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/034,361
; FILING DATE: 24-DEC-1996
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/012,649
; FILING DATE: 01-MAR-1996
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Braham, Susan J.
; REGISTRATION NUMBER: 34,103
; REFERENCE/DOCKET NUMBER: 19603/1062 (D-1906A)
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 716-263-1636
; TELEFAX: 716-263-1600
; INFORMATION FOR SEQ ID NO: 25:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 413 amino acids
; TYPE: amino acid
; STRANDEDNESS: not relevant
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-808-793-25

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Query Match 75.0%; Score 21; DB 2; Length 413;
 Best Local Similarity 100.0%; Pred. No. 5.9e-14;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 94 FIFGSPFTLNLFIVGIIDNF 114

```

RESULT 23
US-07-998-2898-6
; Sequence 6, Application US/079982898
; Patent No. 6027876
; GENERAL INFORMATION:
; APPLICANT: Black, Bruce C
; APPLICANT: Taylor, Martin
; APPLICANT: Heckel, David G
; TITLE OF INVENTION: Method for Monitoring Pesticide
; NUMBER OF SEQUENCES: 40
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Darby & Darby PC
; STREET: 805 Third Avenue
; CITY: New York

```

STATE: New York
COUNTRY: US
ZIP: 10022
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/07/998,289B
FILING DATE: 30-DEC-1992
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Robinson, Joseph R.
REGISTRATION NUMBER: 33,448
REFERENCE/DOCKET NUMBER: 0646/0A939
TELECOMMUNICATION INFORMATION:
TELEPHONE: 212-527-7700
TELEFAX: 212-753-6237
TELEX: 236687
INFORMATION FOR SEQ ID NO: 6:
SEQUENCE CHARACTERISTICS:
LENGTH: 452 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
US-07-998-289B-6

Query Match 75.0%; Score 21; DB 3; Length 452;
Best Local Similarity 100.0%; Pred. No. 6.4e-14;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 FIIFGSFTLNLFIGVLIIDNF 28
DB 126 FIIFGSFTLNLFIGVLIIDNF 146

RESULT 24
US-09-354-147C-7
Sequence 7, Application US/09354147C
Patent No. 6573067
GENERAL INFORMATION:
APPLICANT: Dib-Hajj, Sulayman
TITLE OF INVENTION: Modulation of Sodium Channels in Dorsal Root Ganglia
FILE REFERENCE: 44574-5004-01-US
CURRENT APPLICATION NUMBER: US/09/354,147C
CURRENT FILING DATE: 1999-07-16
PRIOR APPLICATION NUMBER: US 60/072,990
PRIOR FILING DATE: 1998-01-29
PRIOR APPLICATION NUMBER: US 60/109,402
PRIOR FILING DATE: 1998-11-20
PRIOR APPLICATION NUMBER: PCT/US99/02008
PRIOR FILING DATE: 1999-01-29
NUMBER OF SEQ ID NOS: 44
SOFTWARE: Patentin Ver. 2.1
SEQ ID NO 7
LENGTH: 1233
TYPE: PRT
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: UNSURE
LOCATION: (308)
OTHER INFORMATION: Xaa is leu. Xaa results from a "y" in SEQ ID NO: 6.
US-09-354-147C-7

Query Match 75.0%; Score 21; DB 4; Length 1233;
Best Local Similarity 100.0%; Pred. No. 1.5e-13;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 8 FIIFGSFTLNLFIGVLIIDNF 28
DB 1092 FIIFGSFTLNLFIGVLIIDNF 1112

DB 1092 FIIFGSFTLNLFIGVLIIDNF 1112

RESULT 25
US-09-354-147C-8
Sequence 8, Application US/09354147C
Patent No. 6573067
GENERAL INFORMATION:
APPLICANT: Dib-Hajj, Sulayman
TITLE OF INVENTION: Modulation of Sodium Channels in Dorsal Root Ganglia
FILE REFERENCE: 44574-5004-01-US
CURRENT APPLICATION NUMBER: US/09/354,147C
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PRIOR FILING DATE: 1998-11-20
PRIOR APPLICATION NUMBER: PCT/US99/02008
PRIOR FILING DATE: 1999-01-29
NUMBER OF SEQ ID NOS: 44
SOFTWARE: Patentin Ver. 2.1
SEQ ID NO 8
LENGTH: 1243
TYPE: PRT
ORGANISM: Homo sapiens
FEATURE:
OTHER INFORMATION: partial human Nan amino acid seq.
US-09-354-147C-8

Query Match 75.0%; Score 21; DB 4; Length 1243;
Best Local Similarity 100.0%; Pred. No. 1.5e-13;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 FIIFGSFTLNLFIGVLIIDNF 28
DB 1092 FIIFGSFTLNLFIGVLIIDNF 1112

Search completed: January 27, 2005, 17:54:19
Job time : 23.5 secs

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OM protein - protein search, using sw model

Run on: January 27, 2005, 17:32:04 ; Search time 86.5 Seconds

(without alignments)
116.120 Million cell updates/sec

Title: US-10-608-584-29

Perfect score: 28

Sequence: 1 GIFFVSYIIISFLVVMNYAVILENF 28

Scoring table: Gapop 60.0 , Gapext 60.0

Searched: 2002273 seqs, 358729299 residues

Word size : 0

Total number of hits satisfying chosen parameters: 2002273

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Listing first 100 summaries

Database : A_Geneseq_23sep04:*

1: geneeqp19808:*
2: geneeqp19908:*
3: geneeqp20008:*
4: geneeqp20018:*
5: geneeqp20028:*
6: geneeqp20038:*
7: geneeqp20048:*
8: geneeqp20058:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	28	100.0	405	4	AAm5136 Peptide #
2	28	100.0	405	4	ABR34128 Peptide #
3	28	100.0	405	4	AAm27591 Peptide #
4	28	100.0	405	4	ABR28960 Peptide #
5	28	100.0	405	4	ABR19569 Protein #
6	28	100.0	405	4	AAm67299 Human bon
7	28	100.0	405	4	AAm54918 Human bra
8	28	100.0	405	4	AAm56160 Human bra
9	28	100.0	405	4	ABG48961 Human liv
10	28	100.0	405	4	AAm02877 Peptide #
11	28	100.0	405	4	AAm04075 Peptide #
12	28	100.0	405	5	ABG36946 Human pep
13	28	100.0	1855	7	ADBr78597 Human sod
14	28	100.0	1950	7	ADBr78607 Human sod
15	28	100.0	1951	7	ADBr78607 Human sod
16	28	100.0	1951	8	ADBr78607 Human sod
17	28	100.0	1962	5	AAE20511 Human ion
18	28	100.0	1973	5	AAE20516 Human ion
19	28	100.0	1981	7	ABR83185 Human SCN
20	28	100.0	1998	5	AAE20510 Human ion
21	28	100.0	1998	7	ABR83184 Human SCN
22	28	100.0	1999	5	ABR80602 Human sod
23	28	100.0	2000	5	ABR80602 Human sod
24	28	100.0	2000	8	ADK81762 Human Nav
25	28	100.0	2005	4	ABR99676 Human adu

26	28	100.0	2005	4	ABR99677 Human neo
27	28	100.0	2009	5	ABR99674 Human adu
28	28	100.0	2009	5	AAE20515 Human ion
29	28	100.0	2009	5	ABG69292 Human sod
30	28	100.0	2009	5	ABG69291 Human sod
31	28	100.0	2009	5	ABG69293 Human sod
32	28	100.0	2009	5	ABG69289 Human sod
33	28	100.0	2009	5	ABG69289 Human sod
34	28	100.0	2009	5	ABG69290 Human sod
35	28	100.0	2009	5	ABR83180 Human SCN
36	28	100.0	2009	7	ADBr78599 Human sod
37	28	100.0	2009	7	ADBr78599 Human sod
38	28	100.0	2009	7	ADBr78599 Human sod
39	28	100.0	2009	7	ADBr78599 Human sod
40	28	100.0	2009	7	ADBr78599 Human sod
41	28	100.0	2009	7	ADBr78599 Human sod
42	28	100.0	2009	7	ADBr78599 Human sod
43	28	100.0	2009	7	ADBr78599 Human sod
44	28	100.0	2009	7	ADBr78599 Human sod
45	28	100.0	2009	7	ADBr78599 Human sod
46	28	100.0	2009	7	ADBr78599 Human sod
47	28	100.0	2009	7	ADBr78599 Human sod
48	28	100.0	2009	7	ADBr78599 Human sod
49	28	100.0	2009	7	ADBr78599 Human sod
50	28	100.0	2009	7	ADBr78599 Human sod
51	28	100.0	2009	7	ADBr78599 Human sod
52	28	100.0	2009	7	ADBr78599 Human sod
53	28	100.0	2009	7	ADBr78599 Human sod
54	28	100.0	2009	7	ADBr78599 Human sod
55	28	100.0	2009	7	ADBr78599 Human sod
56	28	100.0	2009	7	ADBr78599 Human sod
57	28	100.0	2009	7	ADBr78599 Human sod
58	28	100.0	2009	7	ADBr78599 Human sod
59	28	100.0	2009	7	ADBr78599 Human sod
60	28	100.0	2009	7	ADBr78599 Human sod
61	28	100.0	2009	7	ADBr78599 Human sod
62	28	100.0	2009	7	ADBr78599 Human sod
63	28	100.0	2009	7	ADBr78599 Human sod
64	28	100.0	2009	7	ADBr78599 Human sod
65	28	100.0	2009	7	ADBr78599 Human sod
66	28	100.0	2009	7	ADBr78599 Human sod
67	28	100.0	2009	7	ADBr78599 Human sod
68	28	100.0	2009	7	ADBr78599 Human sod
69	28	100.0	2009	7	ADBr78599 Human sod
70	28	100.0	2009	7	ADBr78599 Human sod
71	28	100.0	2009	7	ADBr78599 Human sod
72	28	100.0	2009	7	ADBr78599 Human sod
73	28	100.0	2009	7	ADBr78599 Human sod
74	28	100.0	2009	7	ADBr78599 Human sod
75	28	100.0	2009	7	ADBr78599 Human sod
76	28	100.0	2009	7	ADBr78599 Human sod
77	28	100.0	2009	7	ADBr78599 Human sod
78	28	100.0	2009	7	ADBr78599 Human sod
79	28	100.0	2009	7	ADBr78599 Human sod
80	28	100.0	2009	7	ADBr78599 Human sod
81	28	100.0	2009	7	ADBr78599 Human sod
82	28	100.0	2009	7	ADBr78599 Human sod
83	28	100.0	2009	7	ADBr78599 Human sod
84	28	100.0	2009	7	ADBr78599 Human sod
85	28	100.0	2009	7	ADBr78599 Human sod
86	28	100.0	2009	7	ADBr78599 Human sod
87	28	100.0	2009	7	ADBr78599 Human sod
88	28	100.0	2009	7	ADBr78599 Human sod
89	28	100.0	2009	7	ADBr78599 Human sod
90	28	100.0	2009	7	ADBr78599 Human sod
91	28	100.0	2009	7	ADBr78599 Human sod
92	28	100.0	2009	7	ADBr78599 Human sod
93	28	100.0	2009	7	ADBr78599 Human sod
94	28	100.0	2009	7	ADBr78599 Human sod
95	28	100.0	2009	7	ADBr78599 Human sod
96	28	100.0	2009	7	ADBr78599 Human sod
97	28	100.0	2009	7	ADBr78599 Human sod
98	28	100.0	2009	7	ADBr78599 Human sod

99 13 46.4 1956 6 ABP72253
100 13 46.4 1956 6 ADA50144

Abp72253 Human PMS
Ada50144 Rat peritp

ALIGNMENTS

RESULT 1
ID AAM15136 standard; protein; 405 AA.
XX AAM15136;

DT 12-OCT-2001 (first entry)

DE Peptide #1570 encoded by probe for measuring cervical gene expression.

KW Probe; human; microarray; gene expression; cervical epithelial cell;

XX Homo sapiens.

PN WO200157278-A2.

PD 09-AUG-2001.

PF 30-JAN-2001; 2001WO-US000670.

PR 04-FEB-2000; 2000US-0180312P.

PR 26-MAY-2000; 2000US-0207456P.

PR 30-JUN-2000; 2000US-00608408.

PR 03-AUG-2000; 2000US-00632366.

PR 21-SEP-2000; 2000US-0234687P.

PR 27-SEP-2000; 2000US-0236359P.

PR 04-OCT-2000; 2000GB-00024263.

XX (MOLE-) MOLECULAR DYNAMICS INC.

XX Penn SG, Hanzel DK, Chen W, Rank DR;

XX WPI; 2001-488901/53.

XX Human genome-derived single exon nucleic acid probes useful for analyzing

PT gene expression in human cervical epithelial cells.

XX Claim 27; SEQ ID NO 19962; 487bp; English.

XX The present invention relates to human single exon nucleic acid probes

CC (SENP; see AAI10068-AA128459). The present sequence is a peptide encoded

CC by one such probe. The SENPs are derived from human HeLa cells. The SENPs

CC can be used to produce a single exon microarray, which can be used for

CC measuring human gene expression in a sample derived from human cervical

CC epithelial cells. By measuring gene expression, the probes are therefore

CC useful in grading and/or staging of diseases of the cervix, notably

CC cervical cancer. Note: The sequence data for this patent did not form

CC part of the printed specification, but was obtained in electronic format

XX directly from WIPO at ftp.wipo.int/pub/published_pct_sequences

SQ Sequence 405 AA;

Query Match 100.0%; Score 28; DB 4; Length 405;
Best Local Similarity 100.0%; Pred. No. 2.6e-20;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GIFFFVSYIIISFLVVMNYIAVILENF 28
DB 158 GIFFFVSYIIISFLVVMNYIAVILENF 185

RESULT 2
ID ABB34128 standard; peptide; 405 AA.
XX ABB34128

AC ABB34128;
XX 04-FEB-2002 (first entry)

DE Peptide #1634 encoded by human foetal liver single exon probe.

KW Human; foetal liver; gene expression; single exon nucleic acid probe.

XX Homo sapiens.

PN WO200157277-A2.

PD 09-AUG-2001.

PF 30-JAN-2001; 2001WO-US000669.

PR 04-FEB-2000; 2000US-0180312P.

PR 26-MAY-2000; 2000US-0207456P.

PR 30-JUN-2000; 2000US-00608408.

PR 03-AUG-2000; 2000US-00632366.

PR 21-SEP-2000; 2000US-0234687P.

PR 27-SEP-2000; 2000US-0236359P.

PR 04-OCT-2000; 2000GB-00024263.

XX (MOLE-) MOLECULAR DYNAMICS INC.

XX Penn SG, Hanzel DK, Chen W, Rank DR;

XX WPI; 2001-483447/52.

XX Human genome-derived single exon nucleic acid probes useful for analyzing

PT gene expression in human fetal liver.

XX Claim 27; SEQ ID NO 26763; 639bp + Sequence listing; English.

XX The invention relates to a single exon nucleic acid probe for measuring

CC human gene expression in a sample derived from human foetal liver. The

CC single exon nucleic acid probes may be used for predicting, measuring and

CC displaying gene expression in samples derived from human fetal liver. The

CC present sequence is a peptide encoded by a single exon nucleic acid probe

CC of the invention. Note: The sequence data for this patent did not form

CC part of the printed specification, but was obtained in electronic format

XX directly from WIPO at ftp.wipo.int/pub/published_pct_sequences

SQ Sequence 405 AA;

Query Match 100.0%; Score 28; DB 4; Length 405;
Best Local Similarity 100.0%; Pred. No. 2.6e-20;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GIFFFVSYIIISFLVVMNYIAVILENF 28
DB 158 GIFFFVSYIIISFLVVMNYIAVILENF 185

RESULT 3
ID AAM27591 standard; protein; 405 AA.
XX AAM27591;
XX 17-OCT-2001 (first entry)

DE Peptide #1628 encoded by probe for measuring placental gene expression.

KW Probe; microarray; human; placenta; antenatal diagnosis;

XX genetic disorder.

XX Homo sapiens.

XX WO200157272-A2.
XX 09-AUG-2001.

XX 30-JAN-2001; 2001WO-US000663.
 PF 04-FEB-2000; 2000US-0180312P.
 PR 26-MAY-2000; 2000US-0207456P.
 PR 30-JUN-2000; 2000US-00608408.
 PR 03-AUG-2000; 2000US-00632366.
 PR 21-SEP-2000; 2000US-0234687P.
 PR 27-SEP-2000; 2000US-0236359P.
 PR 04-OCT-2000; 2000GB-00024263.
 XX (MOLE-) MOLECULAR DYNAMICS INC.
 PA Penn SG, Hanzel DK, Chen W, Rank DR;
 PI WPI; 2001-488897/53.
 DR WPI; 2001-488897/53.
 XX Human genome-derived single exon nucleic acid probes useful for analyzing
 PT gene expression in human placenta.
 PS Claim 27; SEQ ID NO 27860; 654bp; English.
 CC The present invention relates to single exon nucleic acid probes (SENP;
 CC see AI131315-AI157546). The present sequence is a peptide encoded by one
 CC such probe. The probes are useful for producing a microarray for
 CC predicting, measuring and displaying gene expression in samples derived
 CC from human placenta. The probes are useful for antenatal diagnosis of
 CC human genetic disorders
 XX Sequence 405 AA;
 SQ
 Query Match 100.0%; Score 28; DB 4; Length 405;
 Best Local Similarity 100.0%; Pred. No. 2.6e-20;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GIFFFVSYYIIISFLVVMNYIAVILENF 28
 Db 158 GIFFFVSYYIIISFLVVMNYIAVILENF 185

RESULT 4
 ABB28960
 ID ABB28960 standard; peptide; 405 AA.
 XX ABB28960;
 AC
 XX 01-FEB-2002 (first entry)
 DT
 XX Peptide #1611 encoded by breast cell single exon nucleic acid probe.
 DE
 XX Human; microarray; single exon probe; gene expression; breast; disease;
 KW cancer.
 XX Homo sapiens.
 OS
 XX WO200157271-A2.
 PN
 XX 09-AUG-2001.
 PD
 XX 30-JAN-2001; 2001WO-US000662.
 PF
 XX 04-FEB-2000; 2000US-0180312P.
 PR 26-MAY-2000; 2000US-0207456P.
 PR 30-JUN-2000; 2000US-00608408.
 PR 03-AUG-2000; 2000US-00632366.
 PR 21-SEP-2000; 2000US-0234687P.
 PR 27-SEP-2000; 2000US-0236359P.
 PR 04-OCT-2000; 2000GB-00024263.
 XX (MOLE-) MOLECULAR DYNAMICS INC.
 PA Penn SG, Hanzel DK, Chen W, Rank DR;
 PI WPI; 2001-488897/53.
 DR

DR WPI; 2001-496933/54.
 XX New spatially-addressable set of single exon nucleic acid probes, useful
 PT for measuring gene expression in sample derived from human breast,
 PT comprises number of single exon nucleic acid probes.
 PS Claim 27; SEQ ID NO 11928; 327bp + Sequence Listing; English.
 CC The invention relates to a spatially-addressable set of single exon
 CC nucleic acid probes for measuring gene expression in a sample derived
 CC from human breast and BT 474 cells. The method involves contacting the
 CC probes with a collection of detectably labelled nucleic acids derived
 CC from mRNA of human breast, and then measuring the label bound to each
 CC probe of the microarray. The probes are useful for verifying the
 CC expression of regions of genomic DNA predicted to encode proteins. They
 CC are useful for gene discovery, and for determining predisposition and/or
 CC prognosis breast disease. Gene expression analysis is useful for
 CC assessing the toxicity of chemical agents on cells. The microarray of
 CC this invention presents a far greater diversity of probes for measuring
 CC gene expression, with far less bias than expressed sequence tag
 CC microarrays. The method is suitable for rapid production of functional
 CC information from genomic sequence. The present sequence is a peptide
 CC encoded by a single exon nucleic acid probe of the invention. Note: The
 CC sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences
 XX Sequence 405 AA;
 SQ
 Query Match 100.0%; Score 28; DB 4; Length 405;
 Best Local Similarity 100.0%; Pred. No. 2.6e-20;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GIFFFVSYYIIISFLVVMNYIAVILENF 28
 Db 158 GIFFFVSYYIIISFLVVMNYIAVILENF 185

RESULT 5
 ABB19569
 ID ABB19569 standard; protein; 405 AA.
 XX ABB19569;
 AC
 XX 23-JAN-2002 (first entry)
 DT
 XX Protein #1568 encoded by probe for measuring heart cell gene expression.
 DE
 XX Human; gene expression; heart; microarray; vascular system;
 KW cardiovascular disease; hypertension; cardiac arrhythmia;
 KW congenital heart disease.
 XX Homo sapiens.
 OS
 XX WO200157274-A2.
 PN
 XX 09-AUG-2001.
 PD
 XX 30-JAN-2001; 2001WO-US000662.
 PF
 XX 04-FEB-2000; 2000US-0180312P.
 PR 26-MAY-2000; 2000US-0207456P.
 PR 30-JUN-2000; 2000US-00608408.
 PR 03-AUG-2000; 2000US-00632366.
 PR 21-SEP-2000; 2000US-0234687P.
 PR 27-SEP-2000; 2000US-0236359P.
 PR 04-OCT-2000; 2000GB-00024263.
 XX (MOLE-) MOLECULAR DYNAMICS INC.
 PA Penn SG, Hanzel DK, Chen W, Rank DR;
 PI WPI; 2001-488897/53.
 DR

XX Single exon nucleic acid probes for analyzing gene expression in human
 PT hearts.
 XX
 PS Claim 15; SEQ ID NO 21339; 530bp; English.
 XX
 CC The present invention relates to single exon nucleic acid probes for
 CC measuring human gene expression in a sample derived from human heart (see
 CC ABA21535-ABA41305). The present sequence is a protein encoded by one such
 CC probe. The probes may be used for predicting, measuring and displaying
 CC gene expression in samples derived from the human heart via microarrays.
 CC By measuring gene expression, the probes are useful for predicting,
 CC diagnosing, grading, staging, monitoring and prognosing diseases of the
 CC human heart and vascular system e.g. cardiovascular disease.
 CC hypertension, cardiac arrhythmias and congenital heart disease. Note: The
 CC sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences
 CC
 XX
 SQ Sequence 405 AA;

Query Match 100.0%; Score 28; DB 4; Length 405;
 Best Local Similarity 100.0%; Pred. No. 2.6e-20;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 1 GIFFFVSYYIIISFLVVMNYIAVILENF 28
 DB 158 GIFFFVSYYIIISFLVVMNYIAVILENF 185

RESULT 6
 AAM67299
 ID AAM67299 standard; protein; 405 AA.
 XX
 AC AAM67299;
 XX
 DT 06-NOV-2001 (first entry)
 XX
 DE Human bone marrow expressed probe encoded protein SEQ ID NO: 27605.
 XX
 KW Human; bone marrow expressed exon; gene expression analysis; probe;
 KW microarray; cancer; leukaemia; lymphoma; myeloma.
 OS Homo sapiens.
 XX
 PN MO200157276-A2.
 XX
 PD 09-AUG-2001.
 XX
 PF 30-JAN-2001; 2001WO-US000668.
 XX
 PR 04-FEB-2000; 2000US-0180312P.
 PR 26-MAY-2000; 2000US-0207456P.
 PR 30-JUN-2000; 2000US-00608408.
 PR 03-AUG-2000; 2000US-00632366.
 PR 21-SEP-2000; 2000US-0234687P.
 PR 27-SEP-2000; 2000US-0236359P.
 PR 04-OCT-2000; 2000GB-00024263.
 XX
 PA (MOLE-) MOLECULAR DYNAMICS INC.
 XX
 PI Penn SG, Hanzel DK, Chen W, Rank DR;
 XX
 DR WPI; 2001-488900/53.
 XX
 PT Human genome-derived single exon nucleic acid probes useful for analyzing
 PT gene expression in human bone marrow.
 XX
 PS Example 4; SEQ ID NO 27605; 658bp + Sequence Listing; English.
 XX
 CC The present invention provides a number of single exon nucleic acid
 CC probes which are derived from genomic sequences expressed in the human
 CC bone marrow. They can be used to measure gene expression in bone marrow

CC samples, which may enable the improved diagnosis and treatment of cancers
 CC such as lymphoma, leukaemia and myeloma. The present sequence is a
 CC protein encoded by one of the probes of the invention
 CC
 XX
 SQ Sequence 405 AA;

Query Match 100.0%; Score 28; DB 4; Length 405;
 Best Local Similarity 100.0%; Pred. No. 2.6e-20;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 1 GIFFFVSYYIIISFLVVMNYIAVILENF 28
 DB 158 GIFFFVSYYIIISFLVVMNYIAVILENF 185

RESULT 7
 AAM54918
 ID AAM54918 standard; protein; 405 AA.
 XX
 AC AAM54918;
 XX
 DT 05-NOV-2001 (first entry)
 XX
 DE Human brain expressed single exon probe encoded protein SEQ ID NO: 27023.
 XX
 KW Human; brain expressed exon; gene expression analysis; probe; microarray;
 KW Alzheimer's disease; multiple sclerosis; schizophrenia; epilepsy; cancer.
 OS Homo sapiens.
 XX
 PN MO200157275-A2.
 XX
 PD 09-AUG-2001.
 XX
 PF 30-JAN-2001; 2001WO-US000667.
 XX
 PR 04-FEB-2000; 2000US-0180312P.
 PR 26-MAY-2000; 2000US-0207456P.
 PR 30-JUN-2000; 2000US-00608408.
 PR 03-AUG-2000; 2000US-00632366.
 PR 21-SEP-2000; 2000US-0234687P.
 PR 27-SEP-2000; 2000US-0236359P.
 PR 04-OCT-2000; 2000GB-00024263.
 XX
 PA (MOLE-) MOLECULAR DYNAMICS INC.
 XX
 PI Penn SG, Hanzel DK, Chen W, Rank DR;
 XX
 DR WPI; 2001-483446/52.
 XX
 PT Single exon nucleic acid probes for analyzing gene expression in human
 PT brains.
 XX
 PS Example 4; SEQ ID NO 27023; 650bp + Sequence Listing; English.
 XX
 CC The present invention provides a number of single exon nucleic acid
 CC probes which are derived from genomic sequences expressed in the human
 CC brain. They can be used to measure gene expression in brain cell samples,
 CC which may enable the diagnosis and improved treatment of nervous system
 CC diseases such as Alzheimer's disease, multiple sclerosis, schizophrenia,
 CC epilepsy and cancers. The present sequence is a protein encoded by one of
 CC the probes of the invention
 CC
 XX
 SQ Sequence 405 AA;

Query Match 100.0%; Score 28; DB 4; Length 405;
 Best Local Similarity 100.0%; Pred. No. 2.6e-20;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 1 GIFFFVSYYIIISFLVVMNYIAVILENF 28
 DB 158 GIFFFVSYYIIISFLVVMNYIAVILENF 185


```
RESULT 8
AAM56160
ID AAM56160 standard; protein; 405 AA.
XX
AC AAM56160;
XX
DT 05-NOV-2001 (first entry)
XX
DE Human brain expressed single exon probe encoded protein SEQ ID NO: 28265.
XX
KW Human; brain expressed exon; gene expression analysis; probe; microarray;
XX Alzheimer's disease; multiple sclerosis; schizophrenia; epilepsy; cancer.
XX
OS Homo sapiens.
XX
PN WO200157275-A2.
XX
PD 09-AUG-2001.
XX
PF 30-JAN-2001; 2001WO-US000667.
XX
PR 04-FEB-2000; 2000US-0180312P.
PR 26-MAY-2000; 2000US-0207456P.
PR 30-JUN-2000; 2000US-00608408.
PR 03-AUG-2000; 2000US-00632366.
PR 21-SEP-2000; 2000US-0234687P.
PR 27-SEP-2000; 2000US-0236359P.
PR 04-OCT-2000; 2000GB-00024263.
XX
PA (MOLE-) MOLECULAR DYNAMICS INC.
XX
PI Penn SG, Hanzel DK, Chen W, Rank DR;
XX
DR WPI; 2001-483446/52.
XX
PT Single exon nucleic acid probes for analyzing gene expression in human
XX brains.
XX
PS Example 4; SEQ ID NO 28265; 650bp + Sequence Listing; English.
XX
CC The present invention provides a number of single exon nucleic acid
CC probes which are derived from genomic sequences expressed in the human
CC brain. They can be used to measure gene expression in brain cell samples,
CC which may enable the diagnosis and improved treatment of nervous system
CC diseases such as Alzheimer's disease, multiple sclerosis, schizophrenia,
CC epilepsy and cancers. The present sequence is a protein encoded by one of
CC the probes of the invention
XX
SQ Sequence 405 AA;
XX
Query Match 100.0%; Score 28; DB 4; Length 405;
Best Local Similarity 100.0%; Pred. No. 2.6e-20;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GIFFVSYIIISFLVVMNYTAVILENF 28
DB 158 GIFFVSYIIISFLVVMNYTAVILENF 185
XX
RESULT 9
ABG48961
ID ABG48961 standard; peptide; 405 AA.
XX
AC ABG48961;
XX
DT 25-FEB-2003 (first entry)
XX
DE Human liver peptide, SEQ ID No 27609.
XX
KW Human; liver; cirrhosis; hyperlipoproteinaemia; hyperlipidaemia;
XX hypercholesterolaemia; coronary heart disease.
XX
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```
OS Homo sapiens.
XX
PN WO200157273-A2.
XX
PD 09-AUG-2001.
XX
PF 30-JAN-2001; 2001WO-US000664.
XX
PR 04-FEB-2000; 2000US-0180312P.
PR 26-MAY-2000; 2000US-0207456P.
PR 30-JUN-2000; 2000US-00608408.
PR 03-AUG-2000; 2000US-00632366.
PR 21-SEP-2000; 2000US-0234687P.
PR 27-SEP-2000; 2000US-0236359P.
PR 04-OCT-2000; 2000GB-00024263.
XX
PA (MOLE-) MOLECULAR DYNAMICS INC.
XX
PI Penn SG, Hanzel DK, Chen W, Rank DR;
XX
DR WPI; 2001-488898/53.
XX
PT Human genome-derived single exon nucleic acid probes useful for analyzing
XX gene expression in human adult liver.
XX
PS Claim 27; SEQ ID NO 27609; 658bp; English.
XX
CC The invention relates to a single exon nucleic acid probe (SRNP) (1) for
XX measuring human gene expression in a sample derived from human adult
XX liver, comprising one of 13109 defined nucleotide sequences given in the
XX specification (or complements/ fragments). The probe hybridizes at high
XX stringency to a nucleic acid molecule expressed in the human adult liver.
XX (1) may be used for predicting, measuring and displaying gene expression
XX in samples derived from human adult liver. The genes identified may be
XX involved in genetic liver diseases such as cirrhosis,
XX hyperlipoproteinaemia, hyperlipidaemia and hypercholesterolaemia which is
XX associated with coronary heart disease. ABG47348-ABG5930 represent human
XX liver single exon encoded peptides of the invention. Note: The sequence
XX information for this patent does not appear in the printed specification
XX but was obtained in electronic format directly from WIPO at
XX ftp.wipo.int/pub/published_pct_sequences
XX
SQ Sequence 405 AA;
XX
Query Match 100.0%; Score 28; DB 4; Length 405;
Best Local Similarity 100.0%; Pred. No. 2.6e-20;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GIFFVSYIIISFLVVMNYTAVILENF 28
DB 158 GIFFVSYIIISFLVVMNYTAVILENF 185
XX
RESULT 10
AAM02877
ID AAM02877 standard; protein; 405 AA.
XX
AC AAM02877;
XX
DT 09-OCT-2001 (first entry)
XX
DE Peptide #1559 encoded by probe for measuring breast gene expression.
XX
KW Probe; human; breast disease; breast cancer; development disorder;
XX inflammatory disease; proliferative breast disease; non-carcinoma tumour.
XX
OS Homo sapiens.
XX
PN WO200157270-A2.
XX
PD 09-AUG-2001.
XX
PF 29-JAN-2001; 2001WO-US000661.
XX
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XX 04-FEB-2000; 2000US-0180312P.
PR 26-MAY-2000; 2000US-0207456P.
PR 30-JUN-2000; 2000US-0207456P.
PR 03-AUG-2000; 2000US-00608408.
PR 21-SEP-2000; 2000US-00632366.
PR 27-SEP-2000; 2000US-0234687P.
PR 04-OCT-2000; 2000US-0236359P.
XX
XX (MOLE-) MOLECULAR DYNAMICS INC.
XX
XX Penn SG, Hanzel DK, Chen W, Rank DR;
XX WPI; 2001-476286/51.
XX
XX Novel single exon nucleic acid probe used to measuring gene expression in
XX a human breast.
XX
XX Claim 27; SEQ ID NO 11617; 322bp; English.
XX
XX The present invention relates to novel single exon nucleic acid probes
XX (see AAI00010-AA110067). The present sequence is a peptide encoded by one
XX such probe. The probes are useful for measuring human gene expression in
XX a human breast sample, where the probe hybridizes at high stringency to a
XX nucleic acid expressed in the human breast. The probes are useful for
XX predicting, diagnosing, grading, staging, monitoring and prognosing
XX diseases of the human breast, particularly those diseases with polygenic
XX etiology. The diseases include: breast cancer, disorders of development,
XX inflammatory diseases of the breast, fibrocystic changes, proliferative
XX breast disease and non-carcinoma tumours. Note: The sequence data for
XX this patent did not form part of the printed specification, but was
XX obtained in electronic format directly from WIPO at
XX ftp.wipo.int/pub/published_pct_sequences
XX
XX SQ Sequence 405 AA;
XX
XX Query Match 100.0%; Score 28; DB 4; Length 405;
XX Best Local Similarity 100.0%; Pred. No. 2.6e-20;
XX Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 GIFFFVSYYIIISFLVVMNYIAVLENF 28
XX Db 158 GIFFFVSYYIIISFLVVMNYIAVLENF 185
XX
XX RESULT 11
XX ID AAM04075
XX AC AAM04075 standard; protein; 405 AA.
XX
XX DT 09-OCT-2001 (first entry)
XX
XX DE Peptide #2757 encoded by probe for measuring breast gene expression.
XX
XX KW Probe; human; breast disease; breast cancer; development disorder;
XX inflammatory disease; proliferative breast disease; non-carcinoma tumour.
XX
XX OS Homo sapiens.
XX
XX PN WO200157270-A2.
XX
XX PD 09-AUG-2001.
XX
XX PF 29-JAN-2001; 2001WO-US000661.
XX
XX PR 04-FEB-2000; 2000US-0180312P.
XX PR 26-MAY-2000; 2000US-0207456P.
XX PR 30-JUN-2000; 2000US-0207456P.
XX PR 03-AUG-2000; 2000US-00608408.
XX PR 21-SEP-2000; 2000US-00632366.
XX PR 27-SEP-2000; 2000US-0234687P.
XX PR 04-OCT-2000; 2000US-0236359P.
XX

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XX (MOLE-) MOLECULAR DYNAMICS INC.
XX
XX Penn SG, Hanzel DK, Chen W, Rank DR;
XX WPI; 2001-476286/51.
XX
XX Novel single exon nucleic acid probe used to measuring gene expression in
XX a human breast.
XX
XX Claim 27; SEQ ID NO 12815; 322bp; English.
XX
XX The present invention relates to novel single exon nucleic acid probes
XX (see AAI00010-AA110067). The present sequence is a peptide encoded by one
XX such probe. The probes are useful for measuring human gene expression in
XX a human breast sample, where the probe hybridizes at high stringency to a
XX nucleic acid expressed in the human breast. The probes are useful for
XX predicting, diagnosing, grading, staging, monitoring and prognosing
XX diseases of the human breast, particularly those diseases with polygenic
XX etiology. The diseases include: breast cancer, disorders of development,
XX inflammatory diseases of the breast, fibrocystic changes, proliferative
XX breast disease and non-carcinoma tumours. Note: The sequence data for
XX this patent did not form part of the printed specification, but was
XX obtained in electronic format directly from WIPO at
XX ftp.wipo.int/pub/published_pct_sequences
XX
XX SQ Sequence 405 AA;
XX
XX Query Match 100.0%; Score 28; DB 4; Length 405;
XX Best Local Similarity 100.0%; Pred. No. 2.6e-20;
XX Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 GIFFFVSYYIIISFLVVMNYIAVLENF 28
XX Db 158 GIFFFVSYYIIISFLVVMNYIAVLENF 185
XX
XX RESULT 12
XX ID ABG36946
XX AC ABG36946;
XX
XX DT 19-AUG-2002 (first entry)
XX
XX DE Human peptide encoded by genome-derived single exon probe SEQ ID 26611.
XX
XX KW Human; single exon probe; asthma; lung cancer; COPD; ILD;
XX chronic obstructive pulmonary disease; interstitial lung disease;
XX familial idiopathic pulmonary fibrosis; neurofibromatosis;
XX tuberous sclerosis; Gaucher's disease; Niemann-Pick disease;
XX Hereditary-Pudlak syndrome; sarcoidosis; pulmonary haemosiderosis;
XX pulmonary alveolar proteinosis; lymphangioleiomyomatosis; Karsagen syndrome;
XX primary ciliary dyskinesia; fibrocystic pulmonary dysplasia;
XX hyaline membrane disease.
XX
XX OS Homo sapiens.
XX
XX PN WO200186003-A2.
XX
XX PD 15-NOV-2001.
XX
XX PF 30-JAN-2001; 2001WO-US000665.
XX
XX PR 04-FEB-2000; 2000US-0180312P.
XX PR 26-MAY-2000; 2000US-0207456P.
XX PR 30-JUN-2000; 2000US-00608408.
XX PR 03-AUG-2000; 2000US-00632366.
XX PR 21-SEP-2000; 2000US-0234687P.
XX PR 27-SEP-2000; 2000US-0236359P.
XX PR 04-OCT-2000; 2000US-0236359P.
XX

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PA (MOLE-) MOLECULAR DYNAMICS INC.
 XX Penn SG, Hanzel DK, Chen W, Rank DR;
 XX WPI; 2002-114183/15.
 DR Spatially-addressable set of single exon nucleic acid probes, used to
 PT measure gene expression in human lung samples.
 XX
 PS Claim 27; SEQ ID NO 26611; 634pp; English.
 XX
 CC The invention relates to a spatially-addressable set of single exon
 CC nucleic acid probes for measuring gene expression in a sample derived
 CC from human lung comprising single exon nucleic acid probes having one of
 CC 12614 nucleic acid sequences mentioned in the specification, or their
 CC complements or the 12387 open reading frames derived from the 12614
 CC probes. Also included are a microarray comprising the novel set of probes
 CC; the novel set of probes which hybridise at high stringency to a nucleic
 CC acid expressed in the human lung; measuring gene expression in a sample
 CC derived from human lung, comprising (a) contacting the array with a
 CC collection of detectably labeled nucleic acids derived from human lung
 CC mRNA, and (b) measuring the label detectably bound to each probe of the
 CC array; identifying exons in a eukaryotic genome, comprising (a)
 CC algorithmically predicting at least one exon from genomic sequences of
 CC the eukaryote; and (b) detecting specific hybridisation of detectably
 CC labeled nucleic acids from eukaryote lung mRNA, to a single exon probe,
 CC having a fragment identical to the predicted exon, the probe is included
 CC in the above mentioned microarray; assigning exons to a single gene,
 CC comprising (a) identifying exons from genomic sequence by the method
 CC above and (b) measuring the expression of each of the exons in several
 CC tissues and/or cell types using hybridisation to a single exon
 CC microarrays having a probe with the exon, where a common pattern of
 CC expression of the exons in the tissues and/or cell types indicates that
 CC the exons should be assigned to a single gene; a peptide comprising one
 CC of 12011 sequences, mentioned in the specification, or encoded by the
 CC probes/open reading frames (ORF). The probes are used for gene expression
 CC analysis, and for identifying exons in a gene, particularly using human
 CC lung derived mRNA and for the study of lung diseases such as asthma, lung
 CC cancer, chronic obstructive pulmonary disease (COPD), interstitial lung
 CC disease (ILD), familial idiopathic pulmonary fibrosis, neurofibromatosis,
 CC tuberous sclerosis, Gaucher's disease, Niemann-Pick disease, Hermansky-
 CC Pudlak syndrome, sarcoidosis, pulmonary haemosiderosis, pulmonary
 CC histiocytosis, lymphangioleiomyomatosis, pulmonary alveolar proteinosis,
 CC Karaganesi syndrome, fibrocystic pulmonary dysplasia, primary ciliary
 CC dyskinesia, pulmonary hypertension and hyaline membrane disease. The
 CC present sequence is a peptide/protein encoded by a single exon probe of
 CC the invention. Note: The sequence data for this patent did not form part
 CC of the printed specification, but was obtained in electronic format
 CC directly from WIPO at ftp.wipo.int/pub/published_pat_sequences
 XX
 SQ Sequence 405 AA;
 Query Match 100.0%; Score 28; DB 5; Length 405;
 Best Local Similarity 100.0%; Pred. No. 2.6e-20;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GIFFVSYIIISFLVVMYIAVILENF 28
 DB 158 GIFFVSYIIISFLVVMYIAVILENF 185

KW antidepressant; antiparkinsonian; neuroleptic; tranquiliser; analgesic;
 KW neurotropic; antidiabetic; ophthalmological; epilepsy;
 KW ion channel dysfunction; human.
 XX
 OS Synthetic.
 OS Homo sapiens.
 XX
 PN WO2003008574-A1.
 XX
 PD 30-JAN-2003.
 XX
 PF 08-JUL-2002; 2002WO-AU000910.
 XX
 PR 18-JUL-2001; 2001AU-00006452.
 PR 05-MAR-2002; 2002AU-00000910.
 PR 13-MAY-2002; 2002AU-00002292.
 XX
 PA (BION-) BIONOMICS LTD.
 PA (WALL/) WALLACE R W.
 PI Mulley JC, Harkin LA, Dibbens LM, Phillips HA, Heron SE;
 PI Berkovic SF, Scheffer IE;
 DR N-PSDB; ADB78636.
 XX
 DR WPI; 2003-239332/23.
 XX
 PT Identifying predisposition to an ion channel dysfunction, such as
 PT periodic paralysis, cardiac arrhythmias, migraine, Alzheimer's disease,
 PT schizophrenia, anxiety and depression, by detecting encoding-gene
 PT mutation events.
 XX
 PS Claim 13; SEQ ID NO 141; 106pp; English.
 XX
 CC The invention relates to a novel method for identifying a subject
 CC predisposed to a disorder associated with ion channel dysfunction. The
 CC method comprises ascertaining if at least one of the genes encoding ion
 CC channel subunits (ICS) has undergone a mutation event so that a cDNA
 CC derived from the subject has any of 134 nucleotide sequences. The method
 CC of the invention has neurotropic, neuroprotective, inotropic, antipretic,
 CC antiarrhythmic, antidiabetic, antidepressant, antiparkinsonian,
 CC neuroleptic, tranquiliser, analgesic, neurotropic, antidiabetic, and
 CC ophthalmological activity. A polynucleotide of the invention acts as an
 CC ion channel agonist, or ion channel antagonist. The methods, isolated
 CC nucleic acids, polypeptides, antibody, selective agonist, antagonist or
 CC modulator of an ion channel, cells and genetically modified non-human
 CC animal, are useful for the diagnosis and treatment of epilepsy and/or a
 CC disorder associated with ion channel dysfunction, such as hyper- or hypo-
 CC kalemia, periodic paralysis, myotonia, malignant hyperthermia,
 CC myaethenia, cardiac arrhythmias, episodic ataxia, migraine, Alzheimer's
 CC disease, Parkinson's disease, schizophrenia, hyperplexia, anxiety,
 CC depression, phobic obsessive symptoms, neuropathic pain, inflammatory
 CC pain, chronic/acute pain, Bartter's syndrome, polycystic kidney disease,
 CC Dent's disease, hypernatraemic hypoglycaemia of infancy, cystic
 CC fibrosis, congenital stationary night blindness and total colour
 CC blindness. The present sequence represents a mutant protein of the
 CC invention. The sequence data for this patent is not represented in the
 CC printed specification, but was obtained in electronic format directly
 CC from WIPO at ftp.wipo.int/pub/published_pat_sequences.
 XX
 SQ Sequence 1855 AA;
 Query Match 100.0%; Score 28; DB 7; Length 1855;
 Best Local Similarity 100.0%; Pred. No. 1e-19;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GIFFVSYIIISFLVVMYIAVILENF 28
 DB 1762 GIFFVSYIIISFLVVMYIAVILENF 1789

RESULT 14
 ADB78607
 ID ADB78607 standard; protein; 1950 AA.
 DB 1762 GIFFVSYIIISFLVVMYIAVILENF 1789

XX ADB78607;
AC
XX
XX 04-DEC-2003 (first entry)
DT
XX
DE Human sodium channel subunit mutant SEQ ID NO:151.
XX
XX
XX mutein; mutant; ion channel; ion channel subunit; ICS; nootropic;
KM neuroprotective; inotropic; antipruritic; antiarrhythmic; antiallergic;
KM antipruritic; antiparkinsonian; neuroleptic; tranquilizer; analgesic;
KM neuroleptic; antidiabetic; ophthalmological; epilepsy;
XX ion channel dysfunction; human.
XX
XX Synthetic.
OS Homo sapiens.
XX
XX WO2003008574-A1.
PN
XX
XX 30-JAN-2003.
PD
XX
XX 08-JUL-2002; 2002WO-AU000910.
PF
XX
XX 18-JUL-2001; 2001AU-00006452.
PR 05-MAR-2002; 2002AU-00000910.
PR 13-MAY-2002; 2002AU-00002292.
XX
XX (BION-) BIONOMICS LTD.
PA (WALL-) WALLACE R W.
XX
XX Muller JC, Harkin LA, Dibbens LM, Phillips HA, Heron SE;
PI Berkovic SF, Scheffer IE;
XX
XX WPI; 2003-229332/23.
DR N-PSDB; ADB78646.
XX
XX
XX Identifying predisposition to an ion channel dysfunction, such as
PT periodic paralysis, cardiac arrhythmias, migraine, Alzheimer's disease,
PT schizophrenia, anxiety and depression, by detecting encoding-gene
PT mutation events.
XX
XX Claim 13; SEQ ID NO 151; 106pp; English.
XX
XX The invention relates to a novel method for identifying a subject
CC predisposed to a disorder associated with ion channel dysfunction. The
CC method comprises ascertaining if at least one of the genes encoding ion
CC channel subunits (ICS) has undergone a mutation event so that a cDNA
CC derived from the subject has any of 134 nucleotide sequences. The method
CC of the invention has nootropic, neuroprotective, inotropic, antipruritic,
CC antiarrhythmic, antiallergic, antidepressant, antiparkinsonian,
CC neuroleptic, tranquilizer, analgesic, nephroprotective, antiparkinsonian,
CC ophthalmological activity. A polynucleotide of the invention acts as an
CC ion channel agonist, or ion channel antagonist. The methods, isolated
CC nucleic acids, polypeptides, antibody, selective agonist, antagonist or
CC modulator of an ion channel, cells and genetically modified non-human
CC animal, are useful for the diagnosis and treatment of epilepsy and/or a
CC disorder associated with ion channel dysfunction, such as hyper- or hypo-
CC kalemic periodic paralysis, myotonia, malignant hyperthermia, and
CC myasthenia, cardiac arrhythmias, episodic ataxia, migraine, Alzheimer's
CC disease, Parkinson's disease, schizophrenia, hyperkplexia, anxiety,
CC depression, phobic obsessive symptoms, neuropathic pain, inflammatory
CC pain, chronic/acute pain, Bartter's syndrome, polycystic kidney disease,
CC Dent's disease, hyperinsulinaemic hypoglycaemia of infancy, cystic
CC fibrosis, congenital stationary night blindness and total colour
CC blindness. The present sequence represents a mutant protein of the
CC invention. The sequence data for this patent is not represented in the
CC printed specification, but was obtained in electronic format directly
CC from WIPO at ftp.wipo.int/pub/published_pat_sequences.
XX
XX Sequence 1950 AA;

Query Match 100.0%; Score 28; DB 7; Length 1950;
Best Local Similarity 100.0%; Pred. No. 1.1e-19;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GIEFFVSYIIISFLVVMYIAVILENP 28
DB 1697 GIEFFVSYIIISFLVVMYIAVILENP 1724
RESULT 15
ADES9628
ID ADES9628 standard; protein, 1951 AA.
XX
XX ADES9628;
AC
XX
XX 29-JAN-2004 (first entry)
DT
XX
XX Rat Protein NP_037251, SEQ ID NO 5524.
DE
XX
XX Rat; pain; neuronal tissue; gene therapy; spinal segmental nerve injury;
KM chronic constriction injury; CCI; spared nerve injury; SNI; Chung.
XX
XX Rattus norvegicus.
OS
XX
XX WO2003016475-A2.
PN
XX
XX 27-FEB-2003.
PD
XX
XX 14-AUG-2002; 2002WO-US025765.
PF
XX
XX 14-AUG-2001; 2001US-0312147P.
PR 01-NOV-2001; 2001US-0346382P.
PR 26-NOV-2001; 2001US-0333347P.
XX
XX (GENO) GEN HOSPITAL CORP.
PA (FARB) BAYER AG.
XX
XX Woolf C, D'urso D, Befort K, Costigan M;
PI WPI; 2003-268312/26.
DR GENBANK; NP_037251.
XX
XX
XX New composition comprising two or more isolated polypeptides, useful for
PT preparing a medicament for treating pain in an animal.
PT
XX
XX Claim 1; Page; 1017pp; English.
XX
XX The invention discloses a composition comprising two or more isolated rat
CC or human polynucleotides or a polynucleotide which represents a fragment,
CC derivative or allelic variation of the nucleic acid sequence. Also
CC claimed are a vector comprising the novel polynucleotide, a host cell
CC comprising the vector, a method for identifying a nucleotide sequence
CC which is differentially regulated in an animal subjected to pain and a
CC kit to perform the method, an array, a method for identifying an agent
CC that increases or decreases the expression of the polynucleotide sequence
CC subjected to pain, a method for identifying a compound which regulates
CC the expression of a polynucleotide sequence which is differentially
CC expressed in an animal subjected to pain, a method for identifying a
CC compound that regulates the activity of one or more of the
CC polynucleotides, a method for producing a pharmaceutical composition, a
CC method for identifying a compound or small molecule that regulates the
CC activity in an animal of one or more of the polypeptides given in the
CC specification, a method for identifying a compound useful in treating
CC pain and a pharmaceutical composition comprising the one or more
CC polypeptides or their antibodies. The polynucleotide or the compound that
CC modulates its activity is useful for preparing a medicament for treating
CC pain (e.g. spinal segmental nerve injury (Chung), chronic constriction
CC injury (CCI) and spared nerve injury (SNI)) in an animal (e.g. gene
CC therapy). The sequence presented is a rat protein (shown in Table 2 of
CC the specification) which is differentially expressed during pain. Note:
CC The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic form directly from WIPO at
XX ftp.wipo.int/pub/published_pat_sequences.
XX
XX Sequence 1951 AA;

Query Match 100.0%; Score 28; DB 7; Length 1951;
 Best Local Similarity 100.0%; Pred. No. 1.1e-19;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GIFFVSYIIISFLVVMNYIAVILENF 28
 |||||
 DB 1698 GIFFVSYIIISFLVVMNYIAVILENF 1725

RESULT 16
 ADL06576
 ID ADL06576 standard; protein; 1951 AA.

XX AC ADL06576;
 XX DT 20-MAY-2004 (first entry)
 XX DE Human tumour-associated antigenic target (TAT) polypeptide #75.
 XX KW Human; tumour-associated antigenic target; TAT; cell death; tumour;
 KW cancer; cytostatic.

XX OS Homo sapiens.
 XX PN WO2004016225-A2.

XX PD 26-FEB-2004.
 XX PF 19-AUG-2003; 2003WO-US025892.

XX PR 19-AUG-2002; 2002US-0404809P.
 XX PR 21-AUG-2002; 2002US-0405645P.
 XX PR 23-SEP-2002; 2002US-0413192P.
 XX PR 15-OCT-2002; 2002US-0419008P.
 XX PR 15-NOV-2002; 2002US-0426847P.
 XX PR 02-JUL-2003; 2003US-0484959P.

XX PA (GERTH) GENENTECH INC.

XX PI Desauvage FJ, Frantz G, Hillan KJ, Polakis P, Polson A, Smith V;
 PI Spencer SD, Wu TD, Zhang Z;

XX DR WPI; 2004-257144/24.
 XX DR N-PSDB; ADL06499.

XX PT New antibody that binds to a tumor-associated antigenic target (TAT)
 PT polypeptide, useful for preparing a composition for diagnosing or
 PT treating cancer.

XX PS Claim 2; SEQ ID NO 156; 319pp; English.

XX CC The present invention relates to the isolation of human tumour-associated
 CC antigenic target (TAT) polynucleotide and polypeptide sequences. Also
 CC disclosed is an antibody that binds to a TAT polypeptide. The antibody is
 CC a monoclonal antibody, an antibody fragment, a chimeric antibody or a
 CC humanised antibody. It is conjugated to a growth inhibitory agent. It is
 CC produced in bacteria or in CHO cells and induces death of a cell to which
 CC it binds. The antibody is useful for preparing a composition for
 CC diagnosing or treating tumours and cancer. The present sequence
 CC represents a human TAT polypeptide of the invention.

XX SQ Sequence 1951 AA;

Query Match 100.0%; Score 28; DB 8; Length 1951;
 Best Local Similarity 100.0%; Pred. No. 1.1e-19;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GIFFVSYIIISFLVVMNYIAVILENF 28
 |||||
 DB 1698 GIFFVSYIIISFLVVMNYIAVILENF 1725

RESULT 17
 AAE20511
 ID AAE20511 standard; protein; 1962 AA.

XX AC AAE20511;
 XX DT 01-JUL-2002 (first entry)
 XX DE Human ion channel protein #2.

XX KW Human; novel human protein; NHP; voltage-gated sodium channel;
 KW gene therapy; bioreactor; mental disorder; biological disorder;
 KW medical disorder.

XX OS Homo sapiens.

XX FH Key Location/Qualifiers
 FT Misc-difference 981
 FT /note= "Encoded by MTG"
 FT Misc-difference 1056
 FT /note= "Encoded by RCA"

XX PN WO200214498-A2.

XX PD 21-FEB-2002.

XX PF 15-AUG-2001; 2001WO-US025650.

XX PR 16-AUG-2000; 2000US-0225989P.

XX PA (LEXI-) LEXICON GENETICS INC.

XX PI Turner CA, Mathur B, Mathur D;

XX DR WPI; 2002-280757/32.

XX DR N-PSDB; AAD32840.

XX PT Novel polynucleotides encoding human sodium channel proteins,
 PT particularly voltage-gated sodium channel proteins useful for drug
 PT screening, diagnosis and in gene therapy of biological disorders.

XX PS Claim 1; Page 37-41; 83pp; English.

XX CC The present sequence is novel human protein (NHP), ion channel protein.
 CC NHP share structural similarity with mammalian sodium channel proteins
 CC particularly voltage-gated sodium channel proteins. NHP oligonucleotides
 CC are useful as hybridisation probes for screening libraries and assessing
 CC gene expression patterns. Sequences derived from regions adjacent to the
 CC intron/exon boundaries of NHP gene can be used to design primers for use
 CC in amplification assays to detect mutations within the exons, splice
 CC sites, introns that can be used in diagnostics and pharmacogenomics. NHP
 CC nucleotide sequences are useful for drug screening effective in the
 CC treatment of symptomatic or phenotypic manifestations of perturbing the
 CC normal function of NHP in the body, and nucleotide constructs encoding
 CC NHP products are useful to genetically engineer host cells to express NHP
 CC products in vivo. These genetically engineered cells function as
 CC bioreactors in the body delivering a continuous supply of a NHP, a NHP
 CC peptide, or a NHP fusion protein to the body. Nucleotide construct
 CC encoding NHP products are also useful in gene therapy for modulating NHP
 CC expression and to produce genetically engineered host cells to express
 CC NHP products in vivo. NHP nucleotide sequences may also be used as part
 CC of ribozyme and/or triple helix sequences that are useful for NHP gene
 CC regulation. The NHP polypeptides are useful for generating antibodies, as
 CC reagents in diagnostic assays, for identifying other cellular gene
 CC products related to NHP and as reagents in assays for screening for
 CC compounds that are useful in the treatment of mental, biological or
 CC medical disorders and diseases

XX SQ Sequence 1962 AA;

Query Match 100.0%; Score 28; DB 5; Length 1962;
 Best Local Similarity 100.0%; Pred. No. 1.1e-19;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GIFFFVSYYIIISFLVVMNYIAVILENF 28
 DB 1751 GIFFFVSYYIIISFLVVMNYIAVILENF 1778

RESULT 18

ID AAE20516 standard; protein; 1973 AA.

AC AAE20516;

DT 01-JUL-2002 (first entry)

DE Human ion channel protein #7.

KM Human; novel human protein; NHP; voltage-gated sodium channel;
 KW gene therapy; bioreactor; mental disorder; biological disorder;
 XX medical disorder.

OS Homo sapiens.

PH Key Location/Qualifiers

FT Misc-difference 992

FT /note= "Encoded by MTG"

FT Misc-difference 1067

FT /note= "Encoded by RCA"

PN WO200214498-A2.

PD 21-FEB-2002.

PF 15-AUG-2001; 2001WO-US025650.

PR 16-AUG-2000; 2000US-0225989P.

PA (LEXI-) LEXICON GENETICS INC.

PI Turner CA, Mathur B, Mathur D;

DR WPI; 2002-280757/32.

DR N-PSDB; AAD32845.

PT Novel polynucleotides encoding human sodium channel proteins,
 particularly voltage-gated sodium channel proteins useful for drug
 screening, diagnosis and in gene therapy of biological disorders.

PS Claim 1; Page 64-68; 83pp; English.

CC The present sequence is novel human protein (NHP), ion channel protein.
 CC NHP share structural similarity with mammalian sodium channel proteins.
 CC particularly voltage-gated sodium channel proteins. NHP oligonucleotides
 CC are useful as hybridisation probes for screening libraries and assessing
 CC gene expression patterns. Sequences derived from regions adjacent to the
 CC intron/exon boundaries of NHP gene can be used to design primers for use
 CC in amplification assays to detect mutations within the exons, splice
 CC sites, introns that can be used in diagnostics and pharmacogenomics. NHP
 CC nucleotide sequences are useful for drug screening effective in the
 CC treatment of symptomatic or phenotypic manifestations of perturbing the
 CC normal function of NHP in the body, and nucleotide constructs encoding the
 CC NHP products are useful to genetically engineer host cells to express NHP
 CC products *in vivo*. These genetically engineered cells function as
 CC bioreactors in the body delivering a continuous supply of a NHP, a NHP
 CC peptide, or a NHP fusion protein to the body. Nucleotide construct
 CC encoding NHP products are also useful in gene therapy for modulating NHP
 CC expression and to produce genetically engineered host cells to express
 CC NHP products *in vivo*. NHP nucleotide sequences may also be used as part
 CC of ribozyme and/or triple helix sequences that are useful for NHP gene
 CC regulation. The NHP polypeptides are useful for generating antibodies, as
 CC reagents in diagnostic assays, for identifying other cellular gene
 CC products related to NHP and as reagents in assays for screening for
 CC compounds that are useful in the treatment of mental, biological or
 CC medical disorders and diseases

XX SX Sequence 1973 AA;

Query Match 100.0%; Score 28; DB 5; Length 1973;

Best Local Similarity 100.0%; Pred. No. 1.1e-19;

Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GIFFFVSYYIIISFLVVMNYIAVILENF 28
 DB 1762 GIFFFVSYYIIISFLVVMNYIAVILENF 1789

RESULT 19

ID ABR83185 standard; protein; 1981 AA.

AC ABR83185;

DT 15-JAN-2004 (first entry)

DE Human SCN1A splice variant -84P:SCN1ADELP654-681.

KM SCN1A; sodium channel type 1 alpha-subunit; anticonvulsant; analgesic;
 KW neuroprotective; anesthetic; cyrostatic; cerebroprotective; cardiac;
 KW hypotensive; gene therapy; human; splice variant.

OS Homo sapiens.

PN WO2003072751-A2.

PD 04-SEP-2003.

PF 25-FEB-2003; 2003WO-US006010.

PR 25-FEB-2002; 2002US-0359382P.

PA (UYVA-) UNIV VANDERBILT.

PI George AL, Lossin C;

DR WPI; 2003-712725/67.

DR N-PSDB; ACF57880.

PT Recombinantly expressed sodium channel type 1 alpha subunit, useful in
 screening for modulators, for treating e.g. epilepsy.

PS Disclosure; Page 162-169; 176pp; English.

CC The invention relates to a recombinantly expressed and isolated human
 CC SCN1A (sodium channel type 1 alpha-subunit) (I). (II), optionally
 CC incorporated into a cell, is used to screen for specific modulators,
 CC potentially useful as anticonvulsant, antiepileptic, neuroprotective,
 CC analgesic and/or anesthetic agents, e.g. for treating severe myoclonic
 CC epilepsy of infancy, stroke, cardiac arrest, hyperkalemic paralysis,
 CC motor endplate diseases, hypertension, congestive heart failure and
 CC muscular dystrophy also to treat cancer (SCN1A is expressed in prostatic
 CC and metastatic cancer cell lines). These activities can also be provided
 CC by gene therapy vectors that express (I) or the modulators. The
 CC modulators, also antibodies directed against (II), are used to detect
 CC sodium channel polypeptides. The present sequence represents a human
 CC SCN1A splice variant 84P:SCN1ADELP654-681

SQ Sequence 1981 AA;

Query Match 100.0%; Score 28; DB 7; Length 1981;

Best Local Similarity 100.0%; Pred. No. 1.1e-19;

Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GIFFFVSYYIIISFLVVMNYIAVILENF 28
 DB 1734 GIFFFVSYYIIISFLVVMNYIAVILENF 1761

ID	AAE20510	standard; protein; 1998 AA.
AC	AAE20510;	
DT	01-JUL-2002	(first entry)
XX		
DE	Human ion channel protein #1.	
XX		
KW	Human; novel human protein; NHP; voltage-gated sodium channel;	
KW	gene therapy; bioreactor; mental disorder; biological disorder;	
KW	medical disorder.	
XX		
OS	Homo sapiens.	
XX		
FH	Key	Location/Qualifiers
FT	Misc-difference	981
FT	/note= "Encoded by MTC"	
FT	Misc-difference	1056
FT	/note= "Encoded by RCA"	
XX		
PN	WO200214498-A2.	
PD	21-FEB-2002.	
XX		
PF	15-AUG-2001; 2001WO-US025650.	
XX		
PR	16-AUG-2000; 2000US-0225989P.	
XX		
PA	(LEXI-) LEXICON GENETICS INC.	
XX		
P1	Turner CA, Mathur B, Mathur D;	
XX		
DR	WPI; 2002-280757/32.	
DR	N-PSDB; AAD32839.	
XX		
PT	Novel polynucleotides encoding human sodium channel proteins,	
PT	particularly voltage-gated sodium channel proteins useful for drug	
PT	screening, diagnosis and in gene therapy of biological disorders.	
XX		
PS	Claim 1; Page 30-34; 83pp; English.	
XX		
CC	The present sequence is novel human protein (NHP), ion channel protein.	
CC	NHP share structural similarity with mammalian sodium channel proteins	
CC	particularly voltage-gated sodium channel proteins. NHP oligonucleotides	
CC	are useful as hybridisation probes for screening libraries and assessing	
CC	gene expression patterns. Sequences derived from regions adjacent to the	
CC	intron/exon boundaries of NHP gene can be used to design primers for use	
CC	in amplification assays to detect mutations within the exons, splice	
CC	sites, introns that can be used in diagnostics and pharmacogenomics. NHP	
CC	nucleotide sequences are useful for drug screening effective in the	
CC	treatment of symptomatic or phenotypic manifestations of perturbing the	
CC	normal function of NHP in the body, and nucleotide constructs encoding	
CC	NHP products are useful to genetically engineer host cells to express NHP	
CC	products in vivo. These genetically engineered cells function as	
CC	bioreactors in the body delivering a continuous supply of a NHP, a NHP	
CC	peptide, or a NHP fusion protein to the body. Nucleotide construct	
CC	encoding NHP products are also useful in gene therapy for modulating NHP	
CC	expression and to produce genetically engineered host cells to express	
CC	NHP products in vivo. NHP nucleotide sequences may also be used as part	
CC	of ribozyme and/or tripe helix sequences that are useful for NHP gene	
CC	regulation. The NHP polypeptides are useful for generating antibodies, as	
CC	reagents in diagnostic assays, for identifying other cellular gene	
CC	products related to NHP and as reagents in assays for screening for	
CC	compounds that are useful in the treatment of mental, biological or	
CC	medical disorders and diseases	
XX		
XX	Sequence 1998 AA;	
XX		
XX	Query Match	100.0%; Score 28; DB 5; Length 1998;
XX	Best Local Similarity	100.0%; Pred. No. 1.1e-19;
XX	Matches	28; Conservative 0; Mismatches 0; Indels 0; Gaps 0

Oy	1	G F E F V S Y I I S F L V V M M T A V L E N F	28
Dd	1751	G F E F V S Y I I S F L V V M M T A V L E N F	1778
 RESULT 21			
ID	ABR83184	standard; protein; 1998 AA.	
AC	ABR83184;		
DT	15-JAN-2004	(first entry)	
DE	Human SCNA splice variant -33P:SCNADBLP671-681.		
KM	SCNA; sodium channel type 1 alpha-subunit; anticonvulsant; analgesic; neuroprotective; anesthetic; cytostatic; cerebroprotective; cardiac; hypotensive; gene therapy; human; splice variant.		
OS	Homo sapiens.		
PN	WO2003072751-A2.		
PD	04-SEP-2003.		
Pf	25-FEB-2003; 2003MO-US006010.		
PR	25-FEB-2002; 2002US-0359382P.		
PA	(UYVA-) UNIV VANDERBILT.		
PI	George AL, Loshin C;		
DR	WPI; 2003-712725/67.		
DR	N-PSDB; ACF57879.		
PT	Recombinantly expressed sodium channel type 1 alpha subunit, useful in screening for modulators, for treating e.g. epilepsy.		
xx	xx		
P8	Disclosure; Page 148-156; 176pp; English.		
CC	The invention relates to a recombinantly expressed and isolated human SCNA (sodium channel type 1 alpha-subunit) (I). (I), optionally incorporated into a cell, is used to screen for specific modulators, potentially useful as anticonvulsant, antiepileptic, neuroprotective, analgesic and/or anesthetic agents, e.g. for treating severe myoclonic epilepsy of infancy, stroke, cardiac arrest, hyperkalemic paralysis, motor endplate diseases, hypertension, congestive heart failure and muscular dystrophy also to treat cancer (SCNA is expressed in prostatic and metastatic cancer cell lines). These activities can also be provided by gene therapy vectors that express (I) or the modulators. The modulators, also antibodies directed against (I), are used to detect sodium channel polypeptides. The present sequence represents a human SCNA splice variant 33P:SCNADBLP671-681 encoding cDNA		
CC	xx		
CC	xx		
CC	Sequence 1998 AA:		
Query Match	100.0%; Score 28; DB 7; Length 1998;		
Best Local Similarity	100.0%; Pred. No. 1,le-19;		
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0			
Oy	1	G F E F V S Y I I S F L V V M M T A V L E N F	28
Dd	1751	G F E F V S Y I I S F L V V M M T A V L E N F	1778
 RESULT 22			
ID	ABB06026	standard; protein; 1999 AA.	
XX	ABB06026;		
XX	xx		

DT 10-MAY-2002 (first entry)
 XX
 DE Human sodium channel SCN1A protein SEQ ID NO:2.
 XX
 KW Human; sodium channel; SCN1A; chromosome 2q24;
 XX familial hypercalcaemic periodic paralysis; motor endplate disease.
 OS Homo sapiens.
 XX
 XX WO200196552-A1.
 XX
 XX 20-DEC-2001.
 XX
 XX 12-JUN-2001; 2001WO-JP004956.
 XX
 XX 13-JUN-2000; 2000JP-00177540.
 XX 13-JUN-2000; 2000JP-00177544.
 XX
 XX (NISC-) JAPAN SCI & TECHNOLOGY CORP.
 XX
 XX Kanazawa I, Goto J, Jeong S;
 XX
 XX WPI; 2002-098066/13;
 XX N-PSDB; ABL39689.
 XX
 XX Human sodium channels SCN1A and SCN3A and encoded genes, useful in
 PT studying physiological mechanism in which excitant cells participate and
 PT causes of diseases and developing drugs for motor endplate disease.
 XX
 XX Claim 1; Page 40-49; 88pp; Japanese.
 XX
 XX The present invention describes human sodium channels SCN1A and SCN3A.
 CC The present sequence represents the human sodium channel SCN1A. SCN1A and
 CC SCN3A have been located to the human chromosome 2 long arm, positions
 CC 2q24 and 2q24-31 respectively. The sodium channel proteins are useful in
 CC studying the physiological mechanism in which excitant cells participate
 CC and cause human diseases, and in developing remedies for e.g. familial
 CC hypercalcaemic periodic paralysis of extremities and motor endplate
 CC disease
 XX
 XX Sequence 1999 AA;
 SQ
 Query Match 100.0%; Score 28; DB 5; Length 1999;
 Best Local Similarity 100.0%; Pred. No. 1,1e-19;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GIFFFVSYYIIISFLVVMNYIAVILENF 28
 Db 1752 GIFFFVSYYIIISFLVVMNYIAVILENF 1779
 RESULT 23
 ABB06027
 ID ABB06027 standard; protein; 2000 AA.
 XX
 AC ABB06027;
 XX
 DT 10-MAY-2002 (first entry)
 XX
 DE Human sodium channel SCN3A protein SEQ ID NO:4.
 XX
 DE Human; sodium channel; SCN3A; chromosome 2q24-31;
 XX familial hypercalcaemic periodic paralysis; motor endplate disease.
 OS Homo sapiens.
 XX
 XX WO200196552-A1.
 XX
 XX 20-DEC-2001.
 XX
 XX 12-JUN-2001; 2001WO-JP004956.
 XX
 XX 13-JUN-2000; 2000JP-00177540.
 XX

PR 13-JUN-2000; 2000JP-00177544.
 XX
 XX (NISC-) JAPAN SCI & TECHNOLOGY CORP.
 XX
 XX Kanazawa I, Goto J, Jeong S;
 XX
 XX WPI; 2002-098066/13.
 XX N-PSDB; ABL39690.
 XX
 XX Human sodium channels SCN1A and SCN3A and encoded genes, useful in
 PT studying physiological mechanism in which excitant cells participate and
 PT causes of diseases and developing drugs for motor endplate disease.
 XX
 XX Claim 2; Page 72-81; 88pp; Japanese.
 XX
 XX The present invention describes human sodium channels SCN1A and SCN3A.
 CC The present sequence represents the human sodium channel SCN3A. SCN1A and
 CC SCN3A have been located to the human chromosome 2 long arm, positions
 CC 2q24 and 2q24-31 respectively. The sodium channel proteins are useful in
 CC studying the physiological mechanism in which excitant cells participate
 CC and cause human diseases, and in developing remedies for e.g. familial
 CC hypercalcaemic periodic paralysis of extremities and motor endplate
 CC disease
 XX
 XX Sequence 2000 AA;
 SQ
 Query Match 100.0%; Score 28; DB 5; Length 2000;
 Best Local Similarity 100.0%; Pred. No. 1,1e-19;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GIFFFVSYYIIISFLVVMNYIAVILENF 28
 Db 1747 GIFFFVSYYIIISFLVVMNYIAVILENF 1774
 RESULT 24
 ADK81762
 ID ADK81762 standard; protein; 2000 AA.
 XX
 AC ADK81762;
 XX
 DT 20-MAY-2004 (first entry)
 XX
 DE Human Nav1.3 protein.
 XX
 KW Nav1.3; Analgesic; Nootropic; Neuroprotective; post-herpetic neuralgia;
 KW diabetic neuropathy; arthritic pain; migraine headache;
 KW infantile epilepsy; ataxia.
 XX
 OS Homo sapiens.
 XX
 XX WO2004016754-A2.
 XX
 XX 26-FEB-2004.
 XX
 XX 14-AUG-2003; 2003WO-US025465.
 XX
 XX 14-AUG-2002; 2002US-0403416P.
 XX
 XX (PHAA) PHARMACIA CORP.
 XX
 XX Roberds SL;
 XX
 XX WPI; 2004-203785/19.
 XX N-PSDB; ADK81761.
 XX
 XX New antisense compound targeted to a nucleic acid molecule encoding
 PT Nav1.3, useful for treating a disease or condition associated
 PT with Nav1.3, e.g. pain, seizure disorder such as childhood seizure
 PT disorder, or ataxia.
 XX
 XX Disclosure; SEQ ID NO 9096; 417pp; English.
 XX

CC The present invention relates to an antisense compound targeted to a
CC nucleic acid molecule encoding Nav1.3, where the antisense compound
CC specifically hybridizes with and inhibits the expression of Nav1.3. The
CC compound and composition are useful for treating a disease or condition
CC associated with Nav1.3, e.g. pain including but not limited to
CC neuropathic pain, post-herpetic neuralgia, chronic pain, lower back pain,
CC diabetic neuropathy, trigeminal neuropathy, arthritic pain, acute pain,
CC pain from burns, migraine headache, cluster headache, mild-to-moderate
CC headache; seizure disorder such as childhood seizure disorder, including
CC but not limited to neonatal or infantile epilepsy; or ataxia. The present
CC sequence represents human Nav1.3 protein.

XX Sequence 2000 AA;

Query Match 100.0%; Score 28; DB 8; Length 2000;
Best Local Similarity 100.0%; Pred. No. 1.1e-19;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GIFFFVSYYIIISFLVVMNYIAVILENF 28
Db 1747 GIFFFVSYYIIISFLVVMNYIAVILENF 1774

RESULT 25

AAB99676 ID AAB99676 standard; protein; 2005 AA.

XX AAB99676;

XX 04-SEP-2001 (first entry)

XX Human adult form of SCN2A protein sequence SEQ ID NO:35.

XX Human, epilepsy; chromosome 2; SCN1A; SCN2A; SCN3A; identification;
KM diagnosis; mutation; chromosome 2q23-q31; neurological disorder;
KM anticonvulsant; neuroprotective.

XX Homo sapiens.

XX MO200138564-A2.

XX 31-MAY-2001.

XX 24-NOV-2000; 2000MO-CA001404.

XX 26-NOV-1999; 99US-0167623P.

XX (UYMC-) UNIV MCGILL.

XX Rouleau GA, Lafreniere RG, Rochefort D, Cossette P, Ragsdale D;

XX WPI; 2001-355945/37.

XX N-PSDB; AAH55793.

XX Determining a predisposition to epilepsy and/or development of epilepsy
XX comprises determining the genotype of SCN1A, SCN2A and/or SCN3A, or a DNA
XX variant, equivalent, or mutation which shows a linkage disequilibrium.
XX Disclosure; Page 123-130; 268pp; English.

XX The present invention describes a method (M1) of determining an
XX individual's predisposition to epilepsy and/or development of epilepsy,
XX as well as predicting the individual's response to medication. The method
XX comprises determining the genotype of at least one gene selected from
XX SCN1A, SCN2A or SCN3A, or a DNA variant, equivalent, or mutation which
XX shows a linkage disequilibrium. SCN1A, SCN2A and SCN3A are all sodium
XX channel genes located on chromosome 2. The idiopathic generalised
XX epilepsy (IGE) gene is more specifically localised on chromosome 2q23-
XX q31. Compounds identified as modulators of the biological activity of
XX SCN1A, SCN2A or SCN3A proteins or genes, are useful for treating epilepsy
XX or other neurological disorders. They have anticonvulsant and
XX neuroprotective activities. AAH55763 to AAH56164 and AAB99674 to AAB99679
XX represent SCN1A, SCN2A, and SCN3A cDNAs, gene fragments, PCR primers,

CC oligonucleotides and proteins given in the exemplification of the present
CC invention
XX Sequence 2005 AA;

Query Match 100.0%; Score 28; DB 4; Length 2005;
Best Local Similarity 100.0%; Pred. No. 1.1e-19;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GIFFFVSYYIIISFLVVMNYIAVILENF 28
Db 1752 GIFFFVSYYIIISFLVVMNYIAVILENF 1779

Search completed: January 27, 2005, 17:45:18
Job time : 88.5 secs

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OM protein - protein search, using sw model

Run on: January 27, 2005, 17:36:05 ; Search time 17 Seconds
(without alignments)
158.475 Million cell updates/sec

Title: US-10-608-584-29

Perfect score: 28

Sequence: 1 GIFFFVSyllISFLVVMNYAVILENF 28

Scoring table: OLIGO
Gapop 60.0 , Gapext 60.0

Searched: 283416 seqs, 96216763 residues

Word size : 0

Total number of hits satisfying chosen parameters: 283416

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Listing first 100 summaries

Database :
1: p1r1:*
2: p1r2:*
3: p1r3:*
4: p1r4:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	28	100.0	303	2	S29185 sodium channel pro
2	28	100.0	423	2	S29184 sodium channel pro
3	28	100.0	1951	2	S00320 sodium channel pro
4	28	100.0	1983	2	A60054 sodium channel pro
5	28	100.0	2005	2	B25019 sodium channel pro
6	28	100.0	2009	2	A25019 sodium channel pro
7	24	85.7	1977	2	S54771 sodium channel pro
8	20	71.4	70	2	I59194 sodium channel pro
9	17	60.7	1993	2	T30902 sodium channel SCA
10	16	57.1	2005	2	A46269 sodium channel alp
11	14	50.0	324	2	A45752 sodium channel pro
12	14	50.0	1976	2	I56555 sodium channel pro
13	13	46.4	1765	2	T42388 sodium channel alp
14	13	46.4	1957	2	S68453 sodium channel pro
15	12	42.9	1699	2	T31340 voltage-gated sodi
16	11	39.3	1522	2	JC1101 sodium channel pro
17	11	39.3	1595	2	J60084 voltage-gated sodi
18	11	39.3	1739	2	A48298 sodium channel hom
19	11	39.3	1784	2	T43167 sodium channel pro
20	11	39.3	1820	1	CHER sodium channel pro
21	10	35.7	213	2	A30302 sodium channel pro
22	10	35.7	428	2	S35815 sodium channel pro
23	10	35.7	1689	2	S72467 sodium channel pro
24	10	35.7	1820	2	A33299 sodium channel pro
25	10	35.7	2108	2	S72458 sodium channel pro
26	8	28.6	1835	2	I54323 sodium channel alp
27	8	28.6	1836	2	I64893 sodium channel alp
28	8	28.6	1836	2	JS0648 sodium channel alp
29	8	28.6	1836	2	I51964 sodium channel alp

30	8	28.6	1840	1	CHRTM1 sodium channel pro
31	7	25.0	79	2	G82709 hypothetical prote
32	7	25.0	675	2	B75036 hypothetical prote
33	7	25.0	2016	2	A38195 sodium channel pro
34	7	25.0	2019	2	A33996 sodium channel pro
35	7	25.0	2049	2	T43161 sodium channel pro
36	6	21.4	20	2	T26748 hypothetical prote
37	6	21.4	77	2	D64041 hypothetical prote
38	6	21.4	88	2	T17785 hypothetical prote
39	6	21.4	120	2	S62764 NADH2 dehydrogenas
40	6	21.4	122	2	F71180 hypothetical prote
41	6	21.4	161	2	B96537 hypothetical prote
42	6	21.4	171	2	S55958 probable membrane
43	6	21.4	181	2	A87367 transcription regu
44	6	21.4	197	2	D86856 hypothetical prote
45	6	21.4	206	2	B71821 hypothetical prote
46	6	21.4	206	2	T01312 hypothetical prote
47	6	21.4	226	2	G95247 probable heme-bind
48	6	21.4	231	2	S12124 hypothetical prote
49	6	21.4	246	2	D98112 hypothetical prote
50	6	21.4	259	2	A13223 hypothetical prote
51	6	21.4	262	2	S55078 conserved hypothec
52	6	21.4	269	2	F97327 nitroreductase fam
53	6	21.4	274	2	I50682 tenascin - chicken
54	6	21.4	277	2	E75187 sugar abc transpor
55	6	21.4	317	2	B83039 probable permease
56	6	21.4	317	2	H97240 amino acid transpo
57	6	21.4	331	2	H96020 hypothetical prote
58	6	21.4	341	2	T28798 hypothetical prote
59	6	21.4	341	2	E83737 C4-dicarboxylate t
60	6	21.4	370	1	XNVKUD UDPglucose-hexose-
61	6	21.4	371	2	S75778 oligopeptide trans
62	6	21.4	374	2	T25943 hypothetical prote
63	6	21.4	380	2	T11114 ubiquinol-cytochro
64	6	21.4	399	2	T21015 hypothetical prote
65	6	21.4	431	2	T07817 S-locus-specific g
66	6	21.4	439	2	AF1422 cellobiose phospho
67	6	21.4	444	2	I40417 glycerol-3-phospha
68	6	21.4	463	2	E81141 xanthine/uracil pe
69	6	21.4	465	2	C86911 probable cell-divi
70	6	21.4	465	2	T10012 probable cell divi
71	6	21.4	469	2	G70699 probable roxa prot
72	6	21.4	487	2	T27069 hypothetical prote
73	6	21.4	490	2	T33003 hypothetical prote
74	6	21.4	506	2	T23163 hypothetical prote
75	6	21.4	513	2	T03717 GTP-binding protel
76	6	21.4	535	2	AB0146 probable Branched-
77	6	21.4	545	2	S59143 NADH2 dehydrogenas
78	6	21.4	545	2	T01288 protein kinase F27
79	6	21.4	558	2	E70756 hypothetical glyci
80	6	21.4	589	2	H65691 two-component sens
81	6	21.4	627	2	S67257 proline transport
82	6	21.4	634	2	T33528 hypothetical prote
83	6	21.4	666	2	F90069 hypothetical prote
84	6	21.4	710	2	T00055 hypothetical prote
85	6	21.4	713	2	A81317 probable integral
86	6	21.4	862	2	C97343 hypothetical prote
87	6	21.4	939	2	S75908 hypothetical prote
88	6	21.4	1195	2	C87691 hypothetical prote
89	6	21.4	1341	2	T18301 latrophilin-2, sp1
90	6	21.4	1354	2	T18375 latrophilin-2 (sp1
91	6	21.4	1356	2	T18367 latrophilin-2 (sp1
92	6	21.4	1369	2	T18379 latrophilin-2 (sp1
93	6	21.4	1384	2	T18366 latrophilin-2, sp1
94	6	21.4	1397	2	T18377 latrophilin-2 (sp1
95	6	21.4	1399	2	T18370 latrophilin-2 (sp1
96	6	21.4	1407	2	T18381 latrophilin-2 (sp1
97	6	21.4	1412	2	T18380 latrophilin-2 (sp1
98	6	21.4	1420	2	T18385 latrophilin-2 (sp1
99	6	21.4	1422	2	T18383 latrophilin-2, sp1
100	6	21.4	1435	2	T18387 latrophilin-2 (sp1

A:Reference number: A93377; MUID:86146901; PMID:3754035
A:Accession: B25019
A:Molecule type: mRNA
A:Residues: 1-2005 <NOD>
A:Cross-references: UNIPROT:Q63509
A:Experimental source: brain
R:Sarao, R.; Gupta, S.K.; Auid, V.J.; Dunn, R.J.
submitted to the EMBL Data Library, August 1991
A:Description: Developmentally regulated RNA splicing of rat brain sodium channel mRNAs.
A:Reference number: S24803
A:Accession: S24804
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 183-188; 'D', 190-305 <SAR>
A:Cross-references: EMBL:X61149; NID:957074; PIDN:CAA43458.1; PID:957076
C:Superfamily: sodium channel protein
C:Keywords: duplication; ion transport; sodium channel; transmembrane protein; voltage-g

Query Match 100.0%; Score 28; DB 2; Length 2005;
Best Local Similarity 100.0%; Pred. No. 3.2e-20;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GIFFVSYIIISFLVVMNYIAVILENF 28
Db 1752 GIFFVSYIIISFLVVMNYIAVILENF 1779

RESULT 6
A25019
sodium channel protein I - rat
N:Alternate names: sodium channel protein A
C:Species: Rattus norvegicus (Norway rat)
C>Date: 30-Jun-1988 #sequence_revision 30-Jun-1988 #text_change 09-Jul-2004
A:Accession: A25019; S40783; 184764
R:Noda, M.; Ikeda, T.; Kayano, T.; Suzuki, H.; Takeshima, H.; Kurasaki, M.; Takahashi, H.
Nature 340, 188-192, 1986
A>Title: Existence of distinct sodium channel messenger RNAs in rat brain.
A:Reference number: A93377; MUID:86146901; PMID:3754035
A:Accession: A25019
A:Molecule type: mRNA
A:Residues: 1-2009 <NOD>
A:Cross-references: UNIPROT:P04774; GB:X03638; NID:957216; PIDN:CAA27286.1; PID:957217
A:Experimental source: brain
R:Sarao, R.; Gupta, S.K.; Auid, V.J.; Dunn, R.J.
Nucleic Acids Res. 19, 5673-5679, 1991
A>Title: Developmentally regulated alternative RNA splicing of rat brain sodium channel
A:Reference number: S40782; MUID:92051314; PMID:1658739
A:Accession: S40783
A:Molecule type: DNA
A:Residues: 177-253 <SAR>
R:Noda, M.; Numa, S.
J. Recept. Res. 7, 467-497, 1987
A>Title: Structure and function of sodium channel.
A:Reference number: I50536; MUID:87311395; PMID:2442385
A:Accession: I84764
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-2009 <RES>
A:Cross-references: GB:M22253; NID:91041089; PIDN:AAA79965.1; PID:91041089
C:Superfamily: sodium channel protein
C:Keywords: duplication; ion transport; sodium channel; transmembrane protein; voltage-g

Query Match 100.0%; Score 28; DB 2; Length 2009;
Best Local Similarity 100.0%; Pred. No. 3.2e-20;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GIFFVSYIIISFLVVMNYIAVILENF 28
Db 1762 GIFFVSYIIISFLVVMNYIAVILENF 1789

RESULT 7
S54771

sodium channel alpha subunit - human
C:Species: Homo sapiens (man)
C>Date: 27-Oct-1995 #sequence_revision 03-Nov-1995 #text_change 09-Jul-2004
A:Accession: S54771
R:Klugbauer, N.; Lacinova, L.; Flockerzi, V.; Hofmann, F.
EMBO J. 14, 1084-1090, 1995
A>Title: Structure and functional expression of a new member of the tetrodotoxin-sensitive
A:Reference number: S54771; MUID:95237189; PMID:7720699
A:Accession: S54771
A:Status: preliminary; nucleic acid sequence not shown
A:Molecule type: mRNA
A:Residues: 1-1977 <KLU>
A:Cross-references: UNIPROT:Q15858; EMBL:X82835; NID:9758109; PIDN:CAA58042.1; PID:97581
C:Superfamily: sodium channel protein
C:Keywords: duplication

Query Match 85.7%; Score 24; DB 2; Length 1977;
Best Local Similarity 100.0%; Pred. No. 3.4e-16;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 5 FVSYIIISFLVVMNYIAVILENF 28
Db 1729 FVSYIIISFLVVMNYIAVILENF 1752

RESULT 8
I59194
sodium channel protein - human (fragment)
C:Species: Homo sapiens (man)
C>Date: 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change 21-Nov-1997
A:Accession: I59194
R:Han, J.; Lu, C.
Proc. Natl. Acad. Sci. U.S.A. 88, 335-339, 1991
A>Title: Direct amplification of a single dissected chromosomal segment by polymerase ch
A:Reference number: I59194; MUID:9110524; PMID:1846440
A:Accession: I59194
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-70 <RES>
A:Cross-references: GB:M55662; NID:9179560; PID:9553206
C:Superfamily: sodium channel protein
C:Keywords: duplication

Query Match 71.4%; Score 20; DB 2; Length 70;
Best Local Similarity 100.0%; Pred. No. 2.3e-13;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GIFFVSYIIISFLVVMNY 20
Db 51 GIFFVSYIIISFLVVMNY 70

RESULT 9
T30902
sodium channel SCAP1 alpha chain - California sea hare
C:Species: Aplysia californica (California sea hare)
C>Date: 22-Oct-1999 #sequence_revision 22-Oct-1999 #text_change 09-Jul-2004
A:Accession: T30902
R:Dyer, J.R.; Johnston, W.L.; Castellucci, V.F.; Dunn, R.J.
DNA Cell Biol. 16, 347-356, 1997
A>Title: Cloning and tissue distribution of the Aplysia Na+ channel alpha-subunit cDNA.
A:Reference number: Z20929; MUID:97238630; PMID:9115644
A:Accession: T30902
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-1993 <DYB>
A:Cross-references: UNIPROT:P90670; EMBL:U66915; NID:91842248; PID:91842249; PIDN:AA474
C:Superfamily: sodium channel protein

Query Match 60.7%; Score 17; DB 2; Length 1993;
Best Local Similarity 100.0%; Pred. No. 3.7e-09;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 12 SFLVVMNYAVILENF 28
|||
Db 1719 SFLVVMNYAVILENF 1735

RESULT 10

A46269
sodium channel alpha chain HBA - human
C:Species: Homo sapiens (man)
C:Date: 20-Oct-1993 #sequence_revision 18-Nov-1994 #text_change 21-Nov-1997
C:Accession: A46269
R:Ahmed, C.M.; Ware, D.H.; Lee, S.C.; Patten, C.D.; Ferrer-Montiel, A.V.; Schinder, A.F.
Proc. Natl. Acad. Sci. U.S.A. 89, 8220-8224, 1992
A:Title: Primary structure, chromosomal localization, and functional expression of a vol
A:Reference number: A46269; MUID:92390418; PMID:1325650
A:Accession: A46269
A:Molecule type: mRNA
A:Residues: 1-2005 <AM>
A:Cross-references: GB:M94055
A:Experimental source: Brain
A>Note: sequence extracted from NCBI backbone (NCBIP:113082)
C:Genetics:
A:Map position: 2q23-q24.3
C:Superfamily: sodium channel protein
C:Keywords: duplication

Query Match 57.1%; Score 16; DB 2; Length 2005;
Best Local Similarity 100.0%; Pred. No. 3.8e-08;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 GIFFFVSIIISFLV 16
|||
Db 1752 GIFFFVSIIISFLV 1767

RESULT 11

A45752
sodium channel protein PCSC-1 - rat (fragment)
C:Species: Rattus norvegicus (Norway rat)
C:Date: 03-Jun-1993 #sequence_revision 03-Jun-1993 #text_change 09-Jul-2004
C:Accession: A45752
R:Shih, M.N.; Xu, Y.C.; Barrachini, E.; Goodman, R.H.; Cooperman, S.S.; Mandel, G.; Chi
J. Clin. Invest. 84, 331-336, 1989
A:Title: Expression of diverse Na(+) channel messenger RNAs in rat myocardium. Evidence
A:Reference number: A45752; MUID:89292178; PMID:2544627
A:Accession: A45752
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-324 <SL>
A:Cross-references: UNIPROT:Q63360; GB:M27223; NID:g205611; PIDN:AAA1666.1; PID:g205612
C:Superfamily: sodium channel protein
C:Keywords: duplication

Query Match 50.0%; Score 14; DB 2; Length 324;
Best Local Similarity 100.0%; Pred. No. 8.7e-07;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 GIFFFVSIIISFL 14
|||
Db 86 GIFFFVSIIISFL 99

RESULT 12

I56555
sodium channel protein 6 - rat
C:Species: Rattus norvegicus (Norway rat)
C:Date: 26-Jul-1996 #sequence_revision 26-Jul-1996 #text_change 09-Jul-2004
C:Accession: I56555
R:Schaller, K.L.; Krzemien, D.M.; Yarowsky, P.J.; Kueger, B.K.; Caldwell, J.H.
J. Neurosci. 15, 3231-3242, 1995
A:Title: A novel, abundant sodium channel expressed in neurons and glia.
A:Reference number: I56555; MUID:95271284; PMID:7751906
A:Accession: I56555

A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-1976 <RES>
A:Cross-references: UNIPROT:Q63541; GB:LJ39018; NID:g829033; PIDN:AA42059.1; PID:g829034
C:Genetics:
A:Gene: SGP6
C:Superfamily: sodium channel protein
C:Keywords: duplication

Query Match 50.0%; Score 14; DB 2; Length 1976;
Best Local Similarity 100.0%; Pred. No. 3.8e-06;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 GIFFFVSIIISFL 14
|||
Db 1738 GIFFFVSIIISFL 1751

RESULT 13

T42388
sodium channel alpha chain - rat
C:Species: Rattus norvegicus (Norway rat)
C:Date: 03-Dec-1999 #sequence_revision 03-Dec-1999 #text_change 09-Jul-2004
C:Accession: T42388
R:DiB-Hajj, S.D.; Tytrell, L.; Black, J.A.; Waxman, S.G.
Proc. Natl. Acad. Sci. U.S.A. 95, 8963-8968, 1998
A:Title: Nav, a novel voltage-gated Na channel, is expressed preferentially in periph
A:Reference number: 222149; MUID:98338024; PMID:9671787
A:Accession: T42388
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-1765 <DIR>
A:Cross-references: UNIPROT:O88457; EMBL:AF059030; NID:g3372614; PID:g3372615; PIDN:AA4
A:Experimental source: strain Sprague-Dawley; dorsal root ganglia
A>Note: preferentially expressed in sensory neurons within dorsal root ganglia and trige
C:Superfamily: sodium channel protein

Query Match 46.4%; Score 13; DB 2; Length 1765;
Best Local Similarity 100.0%; Pred. No. 3.5e-05;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 16 VVMNYAVILENF 28
|||
Db 1575 VVMNYAVILENF 1587

RESULT 14

S68453
sodium channel protein SNS - rat
C:Species: Rattus norvegicus (Norway rat)
C:Date: 17-Jul-1998 #sequence_revision 17-Jul-1998 #text_change 09-Jul-2004
C:Accession: S68453
R:Akopian, A.N.; Sivikoff, L.; Wood, J.N.
Nature 379, 257-262, 1996
A:Title: A tetrodotoxin-resistant voltage-gated sodium channel expressed by sensory neur
A:Reference number: S68453; MUID:96138382; PMID:8538791
A:Accession: S68453
A:Status: nucleic acid sequence not shown
A:Molecule type: mRNA
A:Residues: 1-1957 <AKO>
A:Cross-references: UNIPROT:Q63554; GB:X92184; NID:g1209466; PIDN:CAA63095.1; PID:g12094
A:Experimental source: dorsal root ganglia
C:Superfamily: sodium channel protein
C:Keywords: sodium channel; transmembrane protein; voltage-gated ion channel
F:132-148/Domain: transmembrane #status predicted <TM1>
F:158-174/Domain: transmembrane #status predicted <TM2>
F:225-241/Domain: transmembrane #status predicted <TM3>
F:249-265/Domain: transmembrane #status predicted <TM4>
F:316-392/Domain: transmembrane #status predicted <TM5>
F:666-682/Domain: transmembrane #status predicted <TM6>
F:702-718/Domain: transmembrane #status predicted <TM7>
F:731-747/Domain: transmembrane #status predicted <TM8>
F:788-804/Domain: transmembrane #status predicted <TM9>

F:865-881/Domain: transmembrane #status predicted <TM10>
 F:1156-1172/Domain: transmembrane #status predicted <TM11>
 F:1194-1210/Domain: transmembrane #status predicted <TM12>
 F:1221-1237/Domain: transmembrane #status predicted <TM13>
 F:1286-1302/Domain: transmembrane #status predicted <TM14>
 F:1400-1416/Domain: transmembrane #status predicted <TM15>
 F:1482-1498/Domain: transmembrane #status predicted <TM16>
 F:1516-1532/Domain: transmembrane #status predicted <TM17>
 F:1566-1562/Domain: transmembrane #status predicted <TM18>
 F:1606-1622/Domain: transmembrane #status predicted <TM19>
 F:1708-1724/Domain: transmembrane #status predicted <TM20>

Query Match 46.4%; Score 13; DB 2; Length 1957;
 Best Local Similarity 100.0%; Pred. No. 3.8e-05;
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 16 VNMVIAVILENF 28
 Db 1715 VNMVIAVILENF 1727

RESULT 15
 T31340
 voltage-gated sodium channel homolog - Bdeiloura candida
 C:Species: Bdeiloura candida
 C:Date: 02-Sep-2000 #sequence_revision 02-Sep-2000 #text_change 09-Jul-2004
 C:Accession: T31340
 R:Jeliorfki, M.C.; Greenberg, R.M.; Anderson, P.A.
 submitted to the EMBL Data Library, March 1997
 A:Description: A putative voltage-gated sodium channel from the turbellarian flatworm Bd
 A:Reference number: 221006
 A:Accession: T31340
 A:Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: mRNA
 A:Residues: 1-1699 <JE2>
 A:Cross-references: UNIPROT:O02037; EMBL:U93074; NID:g1947093; PID:g1947094; PIDN:AAC630
 C:Genetics:
 A:Gene: Na1
 C:Superfamily: sodium channel protein

Query Match 42.9%; Score 12; DB 2; Length 1699;
 Best Local Similarity 100.0%; Pred. No. 0.00035;
 Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 17 VNMVIAVILENF 28
 Db 1536 VNMVIAVILENF 1547

RESULT 16
 JC1101
 sodium channel protein - Bleeker's squid
 C:Species: Loligo bleekeri (Bleeker's squid)
 C:Date: 09-Oct-1992 #sequence_revision 09-Oct-1992 #text_change 09-Jul-2004
 C:Accession: JC1101
 R:Sato, C.; Matsunoto, G.
 Biochem. Biophys. Res. Commun. 186, 61-68, 1992
 A:Title: Primary structure of squid sodium channel deduced from the complementary DNA se
 A:Reference number: JC1101; MUID:92337659; PMID:1339273
 A:Accession: JC1101
 A:Molecule type: mRNA
 A:Residues: 1-1522 <SNT>
 A:Cross-references: UNIPROT:O05973; GB:D14525; NID:G287448; PID:G287449
 C:Superfamily: voltage-dependent calcium channel protein alpha-1 chain
 C:Keywords: phosphoprotein; sodium channel; transmembrane protein; voltage-gated ion cha
 F:51-70/Domain: transmembrane #status predicted <TM1>
 F:78-100/Domain: transmembrane #status predicted <TM2>
 F:113-114/Domain: transmembrane #status predicted <TM3>
 F:114-167/Domain: transmembrane #status predicted <TM4>
 F:180-201/Domain: transmembrane #status predicted <TM5>
 F:309-331/Domain: transmembrane #status predicted <TM6>
 F:407-426/Domain: transmembrane #status predicted <TM7>
 F:443-464/Domain: transmembrane #status predicted <TM8>

F:473-491/Domain: transmembrane #status predicted <TM9>
 F:499-522/Domain: transmembrane #status predicted <TM10>
 F:532-553/Domain: transmembrane #status predicted <TM11>
 F:605-625/Domain: transmembrane #status predicted <TM12>
 F:778-797/Domain: transmembrane #status predicted <TM13>
 F:816-837/Domain: transmembrane #status predicted <TM14>
 F:847-868/Domain: transmembrane #status predicted <TM15>
 F:875-898/Domain: transmembrane #status predicted <TM16>
 F:916-937/Domain: transmembrane #status predicted <TM17>
 F:1010-1032/Domain: transmembrane #status predicted <TM18>
 F:1101-1120/Domain: transmembrane #status predicted <TM19>
 F:1133-1154/Domain: transmembrane #status predicted <TM20>
 F:1163-1184/Domain: transmembrane #status predicted <TM21>
 F:1195-1218/Domain: transmembrane #status predicted <TM22>
 F:1237-1258/Domain: transmembrane #status predicted <TM23>
 F:1324-1346/Domain: transmembrane #status predicted <TM24>

Query Match 39.3%; Score 11; DB 2; Length 1522;
 Best Local Similarity 100.0%; Pred. No. 0.0032;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 18 NMYIAVILENF 28
 Db 1344 NMYIAVILENF 1354

RESULT 17
 JE0084
 voltage-gated sodium channel alpha subunit - hydromedusa (Polyorchis penicillatus)
 N:Alternate names: PpSCN 1
 C:Species: Polyorchis penicillatus
 C:Date: 11-May-1998 #sequence_revision 29-May-1998 #text_change 09-Jul-2004
 A:Accession: JE0084
 R:Spafford, J.D.; Spencer, A.N.; Gallin, W.J.
 Biochem. Biophys. Res. Commun. 244, 772-780, 1998
 A:Title: A putative voltage-gated sodium channel alpha subunit (PpSCN1) from the hydrozo
 A:Reference number: JE0084; MUID:98205797; PMID:9535741
 A:Accession: JE0084
 A:Molecule type: mRNA
 A:Residues: 1-1695 <SPA>
 A:Cross-references: UNIPROT:O62604; GB:AF047380; NID:G3005563; PIDN:AAC38974.1; PID:G300
 C:Comment: This protein is the only pore-forming alpha subunit available to account for
 C:Superfamily: sodium channel protein
 C:Keywords: glycoprotein
 F:201,273,295,584,1065,1082,1089,1428/Binding site: carbohydrate (Asn) (covalent) #statu
 F:201,273,295,584,1065,1082,1089,1428/Binding site: carbohydrate (Asn) (covalent) #statu

Query Match 39.3%; Score 11; DB 2; Length 1695;
 Best Local Similarity 100.0%; Pred. No. 0.0035;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 18 NMYIAVILENF 28
 Db 1463 NMYIAVILENF 1473

RESULT 18
 A48298
 sodium channel homolog - jellyfish (Cyanea capillata)
 C:Species: Cyanea capillata
 C:Date: 19-Mar-1997 #sequence_revision 19-Mar-1997 #text_change 09-Jul-2004
 A:Accession: A48298
 R:Anderson, P.A.V.; Holman, M.A.; Greenberg, R.M.
 Proc. Natl. Acad. Sci. U.S.A. 90, 7419-7423, 1993
 A:Title: Deduced amino acid sequence of a putative sodium channel from the scyphozoan je
 A:Reference number: A48298; MUID:93348284; PMID:8394021
 A:Accession: A48298
 A:Status: preliminary; nucleic acid sequence not shown
 A:Molecule type: mRNA
 A:Residues: 1-1739 <AND>
 A:Cross-references: UNIPROT:Q17314; GB:L15445
 C:Genetics:
 A:Gene: CYN1
 C:Superfamily: sodium channel protein

```

18 NMVIAVILEN 27
   |||||
165 NMVIAVILEN 174

```


RESULT 23

S72467

sodium channel protein para-type alpha chain - German cockroach (strain CSM) (fragment)
C:Species: Blatella germanica (german cockroach)

A:Variety: strain CSM

C:Date: 29-Jul-1997 #sequence_revision 29-Aug-1997 #text_change 09-Jul-2004

C:Accession: S72467; S72487

R:Miyauchi, M.; Ohyama, K.; Dunlap, D.Y.; Matsumura, F.

submitted to the EMBL Data Library, September 1996

A:Description: Cloning and sequencing of the para-type sodium channel gene from susceptible

A:Reference number: S72467

A:Accession: S72467

A:Molecule type: mRNA

A:Residues: 1-1689 <MT>

A:Cross-references: UNIPROT:Q93135; EMBL:U71083; NID:gl633647; PIDN:ABB82037.1; PID:gl63

R:Miyauchi, M.; Ohyama, K.; Dunlap, D.Y.; Matsumura, F.

Mol. Gen. Genet. 252, 61-68, 1996

A:Title: Cloning and sequencing of the para-type sodium channel gene from susceptible an

A:Reference number: S72487; MWID:96397510; PMID:8804404

A:Accession: S72487

A:Molecule type: mRNA

A:Residues: 711-819 <MTW>

A:Cross-references: EMBL:U71083

C:Superfamily: sodium channel protein

C:Keywords: duplication; sodium channel; transmembrane protein

Query Match

Best Local Similarity 35.7%; Score 10; DB 2; Length 1689;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 18 NMVIAVILEN 27

DB 1614 NMVIAVILEN 1623

RESULT 24

A33299

sodium channel protein - fruit fly (Drosophila melanogaster) (fragment)
C:Species: Drosophila melanogaster

C:Date: 20-Dec-1989 #sequence_revision 20-Dec-1989 #text_change 21-Nov-1997

C:Accession: A33299

R:Loughney, K.; Kreber, R.; Ganetzky, B.

Cell 58, 1143-1154, 1989

A:Title: Molecular analysis of the para locus, a sodium channel gene in Drosophila.

A:Reference number: A33299; MWID:89376565; PMID:2550145

A:Accession: A33299

A:Status: preliminary

A:Molecule type: mRNA

A:Residues: 1-1820 <LOU>

A:Cross-references: GB:M32078; GB:M24285

C:Genetics:

A:Gene: FlyBase:Para

A:Cross-references: FlyBase:FBgn0003036

C:Superfamily: sodium channel protein

C:Keywords: duplication; phosphoprotein; transmembrane protein

Query Match

Best Local Similarity 35.7%; Score 10; DB 2; Length 1820;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 18 NMVIAVILEN 27

DB 1809 NMVIAVILEN 1818

RESULT 25

S72458

sodium channel protein para-type alpha chain - house fly (strain Cooper)
C:Species: Musca domestica (house fly)

A:Variety: strain Cooper

C:Date: 24-Oct-1998 #sequence_revision 24-Oct-1998 #text_change 09-Jul-2004

C:Accession: S72458

R:Williamson, M.S.; Martinez-Torres, D.; Hick, C.A.; Devonshire, A.L.

Mol. Gen. Genet. 252, 51-60, 1996

A:Title: Identification of mutations in the housefly para-type sodium channel gene assoc

A:Reference number: S72458; MWID:96397509; PMID:8804403

A:Accession: S72458

A:Molecule type: mRNA

A:Residues: 1-2108 <MTL>

A:Cross-references: UNIPROT:Q94615; EMBL:X96668

A:Experimental source: strain Cooper

C:Genetics:

A:Map position: 3

C:Superfamily: sodium channel protein

C:Keywords: alternative splicing; glycoprotein; phosphoprotein; sodium channel; transmem

F:302,314,332,967,1451,1470/Binding site: carbohydrate (Asn) (covalent) #status predicted

F:541,1208,1582/Binding site: phosphate (Ser) (covalent) #status predicted

Query Match

Best Local Similarity 35.7%; Score 10; DB 2; Length 2108;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 18 NMVIAVILEN 27

DB 1841 NMVIAVILEN 1850

Search completed: January 27, 2005, 17:52:48
Job time : 19 secs

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OM protein - protein search, using sw model

Run on: January 27, 2005, 17:35:20 ; Search time 92.5 Seconds
(without alignments)
174.167 Million cell updates/sec

Title: US-10-608-584-29

Perfect score: 28

Sequence: 1 GIFFVSYITISFLVVMNYAVILENF 28

Scoring table: Gapop 60.0 , Gapext 60.0

Searched: 1825181 seqs, 575374646 residues

Word size: 0

Total number of hits satisfying chosen parameters: 1825181

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Listing first 100 summaries

Database: UniProt 02:*

1: uniprot_sprot:*

2: uniprot_trembl:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	28	100.0	404	2 O8CCS4	O8CCS4 mus musculus
2	28	100.0	510	2 O62242	O62242 mus musculus
3	28	100.0	519	2 O9P2J1	O9P2J1 homo sapien
4	28	100.0	1951	1 C1N3_RAT	P08104 rattus norv
5	28	100.0	1951	2 O9C007	O9C007 homo sapien
6	28	100.0	1981	2 O8IUJ6	O8IUJ6 homo sapien
7	28	100.0	2000	1 C1N3_HUMAN	O9UJ46 homo sapien
8	28	100.0	2005	1 C1N2_HUMAN	O9J250 homo sapien
9	28	100.0	2005	1 C1N2_RAT	P04775 rattus norv
10	28	100.0	2007	2 O9YGN7	O9YGN7 cynops pyrr
11	28	100.0	2009	1 C1N1_HUMAN	P35498 homo sapien
12	28	100.0	2009	1 C1N1_RAT	P04774 rattus norv
13	27	96.4	253	2 O8CNI8	O8CNI8 mus musculus
14	27	85.7	309	2 O62205	O62205 mus musculus
15	24	85.7	1977	2 O15858	O15858 homo sapien
16	24	85.7	1984	2 O28644	O28644 Oryctolagus
17	24	85.7	1984	2 O08562	O08562 rattus norv
18	17	60.7	279	2 O54811	O54811 cavia porce
19	17	60.7	1993	2 P90670	P90670 alysia cal
20	16	57.1	1880	2 O91BFI	O91BFI takifugu pa
21	14	50.0	278	2 O54812	O54812 cavia porce
22	14	50.0	324	2 O63360	O63360 rattus norv
23	14	50.0	364	2 O9P2O6	O9P2O6 homo sapien
24	14	50.0	364	2 O9N2E3	O9N2E3 pongo pygma
25	14	50.0	364	2 O9N2E4	O9N2E4 gorilla gor
26	14	50.0	364	2 O9N2E5	O9N2E5 pan troglod
27	14	50.0	1976	2 O63541	O63541 rattus norv
28	14	50.0	1978	1 C1N8_MOUSE	O9WCU3 mus musculu
29	14	50.0	1978	2 O88420	O88420 rattus norv
30	14	50.0	1980	1 C1N8_HUMAN	O9UCD0 homo sapien
31	14	50.0	1988	2 O88421	O88421 rattus norv

32	13	46.4	523	2 O62243	O62243 mus musculus
33	13	46.4	1765	2 O88457	O88457 rattus norv
34	13	46.4	1765	2 O9JMD4	O9JMD4 mus musculus
35	13	46.4	1765	2 O9R053	O9R053 mus musculus
36	13	46.4	1791	2 O8NDX3	O8NDX3 homo sapien
37	13	46.4	1791	2 O9UHE0	O9UHE0 homo sapien
38	13	46.4	1791	2 O9UIJ3	O9UIJ3 homo sapien
39	13	46.4	1956	2 O9Y5Y9	O9Y5Y9 homo sapien
40	13	46.4	1956	2 O62968	O62968 rattus norv
41	13	46.4	1957	2 O6G1Y3	O6G1Y3 mus musculus
42	13	46.4	1957	2 O63554	O63554 rattus norv
43	13	46.4	1957	2 AAS45602	AAS45602 mus muscu
44	13	46.4	1958	2 P70276	P70276 mus musculu
45	13	46.4	1962	2 O46669	O46669 canis famli
46	12	42.9	46	2 O71A32	O71A32 sus scrofa
47	12	42.9	46	2 AAO11213	AAO11213 sus acroE
48	12	42.9	1599	2 O02037	O02037 bdelioura c
49	11	39.3	1522	1 C1N1_LOBL	C1N1_LOBL
50	11	39.3	1595	2 O62604	O62604 polyorchis
51	11	39.3	1740	2 O17314	O17314 cyanea capi
52	11	39.3	1784	2 O25377	O25377 loligo opal
53	11	39.3	1820	1 C1N4_ELEBL	C1N4_ELEBL
54	10	35.7	428	2 O9TX80	O9TX80 musca domes
55	10	35.7	1089	2 O8IS97	O8IS97 varroa deat
56	10	35.7	1130	2 O9XZC1	O9XZC1 boophilus m
57	10	35.7	1347	2 O7PMT4	O7PMT4 anopheles g
58	10	35.7	1361	2 O7PETH	O7PETH anopheles g
59	10	35.7	1589	2 O9J135	O9J135 blattella g
60	10	35.7	1695	2 O94584	O94584 heliothis v
61	10	35.7	2031	2 O01306	O01306 blattella g
62	10	35.7	2031	2 O01307	O01307 blattella g
63	10	35.7	2051	2 O86D17	O86D17 pediculus h
64	10	35.7	2051	2 O86D18	O86D18 pediculus h
65	10	35.7	2051	2 O86D19	O86D19 pediculus h
66	10	35.7	2058	2 O6DLT4	O6DLT4 aedes albop
67	10	35.7	2064	2 O6DLT3	O6DLT3 aedes aegypt
68	10	35.7	2086	2 O86M38	O86M38 pediculus h
69	10	35.7	2104	2 O25440	O25440 musca domes
70	10	35.7	2105	2 O25439	O25439 musca domes
71	10	35.7	2108	2 O94615	O94615 musca domes
72	10	35.7	2131	1 C1N4_DROME	C1N4_DROME
73	10	35.7	2215	2 O86D77	O86D77 varroa deat
74	8	28.6	487	2 O8AV15	O8AV15 gallus gall
75	8	28.6	1834	2 O28371	O28371 equus cabal
76	8	28.6	1836	1 C1N4_HUMAN	C1N4_HUMAN
77	8	28.6	1840	1 C1N4_RAT	C1N4_RAT
78	8	28.6	1840	2 O70611	O70611 rattus norv
79	8	28.6	1841	2 O9PER0	O9PER0 mus musculus
80	8	25.0	79	2 O9PE09	O9PE09 xyella fas
81	7	25.0	250	2 O96V18	O96V18 sulfolobus
82	7	25.0	397	2 O7ZVJ1	O7ZVJ1 brachydanio
83	7	25.0	444	2 O7ZVU7	O7ZVU7 leprospira
84	7	25.0	444	2 O8FP94	O8FP94 leprospira
85	7	25.0	444	2 AAS68827	AAS68827 leprospira
86	7	25.0	616	2 O7QWH6	O7QWH6 giardia lam
87	7	25.0	640	1 PTMA_BUCBP	PTMA_BUCBP
88	7	25.0	649	2 O7ZSK6	O7ZSK6 leprospira
89	7	25.0	649	2 O8F303	O8F303 leprospira
90	7	25.0	649	2 AAS69976	AAS69976 leprospira
91	7	25.0	656	2 O81260	O81260 plasmodium
92	7	25.0	662	2 O8TU41	O8TU41 methanosarc
93	7	25.0	678	2 O9UJ26	O9UJ26 pyrococcus
94	7	25.0	678	2 O6HCZ6	O6HCZ6 bacillus th
95	7	25.0	678	2 O81L64	O81L64 bacillus an
96	7	25.0	678	2 AAT33890	AAT33890 bacillus
97	7	25.0	1394	2 O8H8L9	O8H8L9 oryza sativ
98	7	25.0	1717	2 O90519	O90519 fuqua rubrip
99	7	25.0	1949	2 O9DF53	O9DF53 brachydanio
100	7	25.0	1962	2 O75RX9	O75RX9 homo sapien

ALIGNMENTS

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RESULT 1
ID Q8CCS4 PRELIMINARY; PRT; 404 AA.
AC O8CCS4
DT 01-MAR-2003 (TrEMBLrel. 23, Created)
DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE Mus musculus adult male olfactory brain cDNA, RIKEN full-length
DE enriched library, clone:643040810 product:sodium channel protein II
DE homolog (Fragment).
GN Name:A23005E3R1k.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Olfactory brain;
RX MEDLINE=99279253; PubMed=10349636;
RA Carninci P., Hayashizaki Y.;
RT "High-efficiency full-length cDNA cloning.";
RL Meth. Enzymol. 303:19-44 (1999).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Olfactory brain;
RX MEDLINE=21085660; PubMed=11217851;
RA RIKEN PANTOM Consortium;
RT "Functional annotation of a full-length mouse cDNA collection.";
RL Nature 409:685-690 (2001).
RN [3]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Olfactory brain;
RA The FANTOM Consortium;
RT "Analysis of the mouse transcriptome based on functional annotation of
RT 60,770 full-length cDNAs.";
RL Nature 420:563-573 (2002).
RN [4]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Olfactory brain;
RX MEDLINE=20499374; PubMed=11042159;
RA Carninci P., Shibata Y., Hayata N., Sugahara Y., Shibata K., Itoh M.,
RA Kono H., Okazaki Y., Muramatsu M., Hayashizaki Y.;
RT "Normalization and subtraction of cap-trapper-selected cDNAs to
RT prepare full-length cDNA libraries for rapid discovery of new genes.";
RL Genome Res. 10:1617-1630 (2000).
RN [5]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Olfactory brain;
RX MEDLINE=20530913; PubMed=11076861;
RA Shibata K., Itoh M., Aizawa K., Nagaoka S., Sasaki N., Carninci P.,
RA Kono H., Akiyama J., Nishi K., Kitsuai T., Tachiro H., Itoh M.,
RA Sumi N., Ishii Y., Nakamura S., Harada M., Nishino T., Harada A.,
RA Yamamoto R., Matsuno H., Sakaguchi S., Ikegami T., Kasahagi K.,
RA Fujiwaka S., Inoue K., Togawa Y., Izawa M., Ohara E., Watabiki M.,
RA Yoneda Y., Ishikawa T., Ozawa K., Tanaka T., Matsura S., Kawai J.,
RA Okazaki Y., Muramatsu M., Inoue Y., Kira A., Hayashizaki Y.;
RT "RIKEN integrated sequence analysis (RISA) system-384 format
RT sequencing pipeline with 384 multicapillary sequencer.";
RL Genome Res. 10:1757-1771 (2000).
RN [6]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Olfactory brain;
RA Adachi U., Aizawa K., Akiyama T., Aizawa T., Bono H., Carninci P.,
RA Fukuda S., Furuno M., Hanagaki T., Hara A., Hashizume W.,
RA Hayashida K., Hayatsu N., Hiramoto K., Hiroka T., Hirozane T.,
RA Hori F., Imotani K., Ishii Y., Itoh M., Kagawa T., Kasukawa T.,
RA Katoh H., Kawai J., Kojima Y., Kondo S., Kono H., Kouda M., Koya S.,
RA Kurihara C., Matsuyama T., Miyazaki A., Murata M., Nakamura M.,
RA Nihei K., Nomura K., Numazaki R., Ohno M., Ohsato N., Okazaki Y.,
RA Saito R., Saitoh H., Sakai C., Sakai K., Sakazume N., Sano H.,
RA Sasaki D., Shibata K., Shinagawa A., Shitaki T., Sogabe Y., Tagami M.,

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RA Tagawa A., Takahashi F., Takaku-Akahira S., Takeda Y., Tanaka T.,
RA Tomaru A., Toya T., Yasunishi A., Muramatsu M., Hayashizaki Y.;
RL Submitted (JUL-2001) to the EMBL/GenBank/DBJ databases.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
CC -1- SIMILARITY: Belongs to the sodium channel family.
DR EMBL; AK032187; BAC27748.1; -.
DR MGD; MG1:2444703; A23005E3R1k.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0001518; C:voltage-gated sodium channel complex; IEA.
DR GO; GO:0005261; F:cation channel activity; IEA.
DR GO; GO:0006812; F:cation transport; IEA.
DR GO; GO:0006814; P:sodium ion transport; IEA.
DR GO; GO:0006812; P:cation transport; IEA.
DR InterPro; IPR001682; Ca/Na pore.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IQ_region.
DR InterPro; IPR005820; M-channel_nlg.
DR InterPro; IPR001696; Na_channel.
DR Pfam; PF00612; IQ_1.
DR SMART; SM00015; IQ_1.
DR PROSITE; PS00096; IQ_1.
KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
KW Transport; Voltage-gated channel.
FT NON_TER
SQ SEQUENCE 404 AA; 45671 MW; C5ED26B0F080C09 CRC64;
Query Match 100.0%; Score 28; DB 2; Length 404;
Best Local Similarity 100.0%; Pred. No. 28-19;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GIFFVSYYIIISFLVVMYIAVLENF 28
Db 151 GIFFVSYYIIISFLVVMYIAVLENF 178
RESULT 2
ID Q62242 PRELIMINARY; PRT; 510 AA.
AC Q62242
DT 01-NOV-1996 (TrEMBLrel. 01, Created)
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE Sodium channel (Fragment).
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Brain;
RA Jover E., Shah V.;
RL Submitted (JUN-1995) to the EMBL/GenBank/DBJ databases.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
CC -1- SIMILARITY: Belongs to the sodium channel family.
DR EMBL; L42341; AA67695.1; -.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0001518; C:voltage-gated sodium channel complex; IEA.
DR GO; GO:0005261; F:cation channel activity; IEA.
DR GO; GO:0006812; F:cation-gated sodium channel activity; IEA.
DR GO; GO:0006814; P:sodium ion transport; IEA.
DR InterPro; IPR001682; Ca/Na pore.
DR InterPro; IPR002111; Cat_channel_TrypL.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IQ_region.
DR InterPro; IPR005820; M-channel_nlg.
DR InterPro; IPR001696; Na_channel.
DR Pfam; PF00520; Ion_trans; 1.
DR Pfam; PF00612; IQ_1.
DR PRINTS; PR00170; NACHANNEL.

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DR SMART; SM00015; IQ; 1.
 DR PROSITE; PSS0096; IQ; 1.
 KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
 FT NON_TER 510 510
 SQ SEQUENCE 510 AA; 58397 MW; 02DCC7DAED3796E8 CRC64;
 Query Match 100.0%; Score 28; DB 2; Length 510;
 Best Local Similarity 100.0%; Pred. No. 2.4e-19;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GIFFFVSYYIIISFLVVMNYIAVILENF 28
 |||||
 Db 304 GIFFFVSYYIIISFLVVMNYIAVILENF 331

RESULT 3
 QP2J1 PRELIMINARY; PRT; 519 AA.
 AC QP2J1;
 DT 01-OCT-2000 (TrEMBLrel. 15, Created)
 DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
 DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
 DE KIAA1356 protein (Fragment).
 GN Name=KIAA1356;
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Brain;
 RX MEDLINE=20181126; PubMed=10718198; Hirosewa M., Ohara O.;
 RA Nagase T., Kikuno R., Ishikawa K., Hirosewa M., Ohara O.;
 RT "Prediction of the coding sequences of unidentified human genes. XVI.
 RT The complete sequences of 150 new cDNA clones from brain which code
 RT for large proteins in vitro.";
 RL DNA Res. 7:65-73(2000).
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
 CC -1- SIMILARITY: Belongs to the sodium channel family.
 DR EMBL; AB037777; BAA92594.1; -;
 DR GO; GO:0016021; C:integral to membrane; IEA.
 DR GO; GO:0005151; C:voltage-gated sodium channel complex; IEA.
 DR GO; GO:0005261; P:cation channel activity; IEA.
 DR GO; GO:0005248; F:voltage-gated sodium channel activity; IEA.
 DR GO; GO:0006814; P:sodium ion transport; IEA.
 DR GO; GO:0006814; P:sodium ion transport; IEA.
 DR InterPro; IPR002111; Cat_channel_TrpL.
 DR InterPro; IPR005821; Ion_trans.
 DR InterPro; IPR000048; IQ_region.
 DR InterPro; IPR005820; M+channel_nlg.
 DR InterPro; IPR001696; Na_channel.
 DR Pfam; PF00520; Ion_trans. 1.
 DR Pfam; PF00612; IQ. 1.
 DR PRINTS; PR00170; NACHANNEL.
 DR SMART; SM00015; IQ; 1.
 DR PROSITE; PSS0096; IQ; 1.
 KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
 KW Transport; Voltage-gated channel.
 FT NON_TER 1 1
 SQ SEQUENCE 519 AA; 58982 MW; 3E8F7ED5E26835FC CRC64;

CIN3 RAT
 ID CIN3 RAT STANDARD; PRT; 1951 AA.
 AC P08104;
 DT 01-AUG-1988 (Rel. 08, Created)
 DT 01-AUG-1988 (Rel. 08, Last sequence update)
 DT 01-OCT-2004 (Rel. 45, Last annotation update)
 DE Sodium channel protein type III alpha subunit (Voltage-gated sodium
 DE channel alpha subunit Nav1.3) (Sodium channel protein, brain III alpha
 DE subunit) (Voltage-gated sodium channel subtype III).
 GN Name=Scn3a;
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 OX NCBI_TaxID=10116;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=Miscar;
 RX MEDLINE=88137594; PubMed=2449363;
 RA Kayano T., Noda M., Flockerzi V., Takahashi H., Numa S.;
 RT "Primary structure of rat brain sodium channel III deduced from the
 RT cDNA sequence.";
 RL FEBS Lett. 228:187-194(1988).
 CC -1- FUNCTION: Mediates the voltage-dependent sodium ion permeability
 CC of excitable membranes. Assuming opened or closed conformations in
 CC response to the voltage difference across the membrane, the
 CC protein forms a sodium-selective channel through which Na(+) ions
 CC may pass in accordance with their electrochemical gradient.
 CC -1- SUBUNIT: The sodium channel consists of a large polypeptide and 2-
 CC 3 smaller ones. This sequence represents a large polypeptide.
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
 CC -1- DOMAIN: The sequence contains 4 internal repeats, each with 5
 CC hydrophobic segments (S1,S2,S3,S5,S6) and one positively charged
 CC segment (S4). Segments S4 are probably the voltage-sensors and are
 CC characterized by a series of positively charged amino acids at
 CC every third position.
 CC -1- SIMILARITY: Belongs to the sodium channel family.
 CC -1- SIMILARITY: Contains 1 IQ domain.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
 CC the European Bioinformatics Institute. There are no restrictions on its
 CC use by non-profit institutions as long as its content is in no way
 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@sib-sib.ch).
 CC -----
 DR EMBL; Y00766; CAA68735.1; -;
 DR PIR; S00320; S00320.
 DR PDB; 1OG9; NMR; A=156-176.
 DR RGD; 3635; Scn3a.
 DR InterPro; IPR001682; Ca/Na_pore.
 DR InterPro; IPR002111; Cat_channel_TrpL.
 DR InterPro; IPR005821; Ion_trans.
 DR InterPro; IPR000048; IQ_region.
 DR InterPro; IPR005820; M+channel_nlg.
 DR InterPro; IPR001696; Na_channel.
 DR InterPro; IPR010526; Na_trans_assoc.
 DR Pfam; PF00520; Ion_trans. 4.
 DR Pfam; PF00612; IQ. 1.
 DR Pfam; PF06512; Na_trans_assoc. 1.
 DR PRINTS; PR00170; NACHANNEL.
 DR PROSITE; PSS0096; IQ; FALSE NEG.
 KW 3D-structure; Glycoprotein; Ion transport; Ionic channel;
 KW Multigene family; Repeat; Sodium channel; Transmembrane;
 KW Voltage-gated channel.
 FT REPEAT 110 455 1.
 FT REPEAT 693 965 II.
 FT REPEAT 1139 1450 III.
 FT REPEAT 1459 1757 IV.
 FT TRANSMEM 124 147 S1 of repeat I.
 FT TRANSMEM 156 175 S2 of repeat I.
 FT TRANSMEM 189 207 S3 of repeat I.
 FT TRANSMEM 214 233 S4 of repeat I.

RESULT 4

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FT TRANSMEM 249 273 S5 of repeat I.
FT TRANSMEM 401 426 S6 of repeat I.
FT TRANSMEM 706 730 S1 of repeat II.
FT TRANSMEM 742 765 S2 of repeat II.
FT TRANSMEM 774 793 S3 of repeat II.
FT TRANSMEM 800 820 S4 of repeat II.
FT TRANSMEM 836 856 S5 of repeat II.
FT TRANSMEM 910 935 S6 of repeat II.
FT TRANSMEM 1153 1176 S1 of repeat III.
FT TRANSMEM 1190 1215 S2 of repeat III.
FT TRANSMEM 1222 1243 S3 of repeat III.
FT TRANSMEM 1248 1269 S4 of repeat III.
FT TRANSMEM 1289 1310 S5 of repeat III.
FT TRANSMEM 1393 1419 S6 of repeat III.
FT TRANSMEM 1473 1496 S1 of repeat IV.
FT TRANSMEM 1508 1531 S2 of repeat IV.
FT TRANSMEM 1538 1561 S3 of repeat IV.
FT TRANSMEM 1572 1593 S4 of repeat IV.
FT TRANSMEM 1609 1631 S5 of repeat IV.
FT TRANSMEM 1698 1722 S6 of repeat IV.
FT CARBOHYD 211 211 N-1linked (GlcNAc. . .) (Potential).
FT CARBOHYD 290 290 N-1linked (GlcNAc. . .) (Potential).
FT CARBOHYD 296 296 N-1linked (GlcNAc. . .) (Potential).
FT CARBOHYD 302 302 N-1linked (GlcNAc. . .) (Potential).
FT CARBOHYD 307 307 N-1linked (GlcNAc. . .) (Potential).
FT CARBOHYD 339 339 N-1linked (GlcNAc. . .) (Potential).
FT CARBOHYD 424 424 N-1linked (GlcNAc. . .) (Potential).
FT CARBOHYD 835 835 N-1linked (GlcNAc. . .) (Potential).
FT CARBOHYD 1002 1002 N-1linked (GlcNAc. . .) (Potential).
FT CARBOHYD 1019 1019 N-1linked (GlcNAc. . .) (Potential).
FT CARBOHYD 1085 1085 N-1linked (GlcNAc. . .) (Potential).
FT CARBOHYD 1317 1317 N-1linked (GlcNAc. . .) (Potential).
FT CARBOHYD 1331 1331 N-1linked (GlcNAc. . .) (Potential).
SQ SEQUENCE 1951 AA; 22135 MW; 745E851524BD10E CRC64;

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Query Match 100.0%; Score 28; DB 1; Length 1951;
Best local Similarity 100.0%; Pred. No. 6.5e-19;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GIFFVSYIIISFLVVMYIAVLLENF 28
DB 1698 GIFFVSYIIISFLVVMYIAVLLENF 1725

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RESULT 5
Q9C007 PRELIMINARY; PRT; 1951 AA.
AC Q9C007;
DT 01-JUN-2001 (TREMBlrel. 17, Created)
DT 01-JUN-2001 (TREMBlrel. 17, last sequence update)
DE 01-MAR-2004 (TREMBlrel. 26, last annotation update)
DE Voltage-gated sodium channel alpha subunit splice variant SCN3A-B.
GN Name=SCN3A;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Jeong S.-Y., Goto J., Kanazawa I.;
RC Submitted (JAN-2000) to the EMBL/GenBank/DBJ databases.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
CC -1- SIMILARITY: Belongs to the sodium channel family.
DR EMBL; AF225986; AAK00218.1; -.
DR HSSP; P04775; 1BY1.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:001518; C:voltage-gated sodium channel complex; IEA.
DR GO; GO:0005261; P:cation channel activity; IEA.
DR GO; GO:0005248; P:voltage-gated sodium channel activity; IEA.
DR GO; GO:0006814; P:cation transport; IEA.
DR GO; GO:0006814; P:sodium ion transport; IEA.
DR InterPro; IPR001682; Ca/Na_pore.

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DR InterPro; IPR002111; Cat_channel_TrpL.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IQ_Region.
DR InterPro; IPR005820; M_channel_nlg.
DR InterPro; IPR001696; Na_channel.
DR InterPro; IPR010526; Na_trans_assoc.
DR Pfam; PF00520; Ion_trans_4.
DR Pfam; PF06512; IQ_1.
DR Pfam; PF06512; Na_trans_assoc. 1.
DR PRINTS; PR00170; NACHANNEL.
DR SMART; SM00015; IQ_1.
DR PROSITE; PS50096; IQ_1.
KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
KW transport; Voltage-gated channel.
SQ SEQUENCE 1951 AA; 221517 MW; 99AD4C032CE124AB CRC64;

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Query Match 100.0%; Score 28; DB 2; Length 1951;
Best local Similarity 100.0%; Pred. No. 6.5e-19;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GIFFVSYIIISFLVVMYIAVLLENF 28
DB 1698 GIFFVSYIIISFLVVMYIAVLLENF 1725

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RESULT 6
Q8IUJ6 PRELIMINARY; PRT; 1981 AA.
AC Q8IUJ6;
DT 01-MAR-2003 (TREMBlrel. 23, Created)
DT 01-MAR-2003 (TREMBlrel. 23, last sequence update)
DE 01-MAR-2004 (TREMBlrel. 26, last annotation update)
DE Voltage-gated sodium channel alpha 1 subunit.
GN Name=SCN1A;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Tissue-Normal Brain;
RA Ouchida M., Ohmori I.;
RC Submitted (DEC-2002) to the EMBL/GenBank/DBJ databases.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
CC -1- SIMILARITY: Belongs to the sodium channel family.
DR EMBL; AB098335; BA045228.1; -.
DR HSSP; P04775; 1BY1.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:000518; C:voltage-gated sodium channel complex; IEA.
DR GO; GO:0005261; P:cation channel activity; IEA.
DR GO; GO:0005248; P:voltage-gated sodium channel activity; IEA.
DR GO; GO:0006812; P:cation transport; IEA.
DR GO; GO:0006814; P:sodium ion transport; IEA.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR002111; Cat_channel_TrpL.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IQ_Region.
DR InterPro; IPR005820; M_channel_nlg.
DR InterPro; IPR001696; Na_channel.
DR InterPro; IPR008051; Na_channel1.
DR InterPro; IPR010526; Na_trans_assoc.
DR InterPro; IPR000100; Ribonuclease_P.
DR Pfam; PF00520; Ion_trans_4.
DR Pfam; PF06512; IQ_1.
DR Pfam; PF06512; Na_trans_assoc. 1.
DR PRINTS; PR00170; NACHANNEL.
DR PRINTS; PR01664; NACHANNEL1.
DR SMART; SM00015; IQ_1.
DR PROSITE; PS00648; RIBONUCLEASE_P; UNKNOWN 1.
KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
KW transport; Voltage-gated channel.
SQ SEQUENCE 1981 AA; 226201 MW; B1D6946D491B7AD CRC64;

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Query Match 100.0%; Score 28; DB 2; Length 1981;
Best Local Similarity 100.0%; Pred. No. 6,6e-19;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GIFFFVSIIISPLVVNNYIVILNPF 28
DB 1734 GIFFFVSIIISPLVVNNYIVILNPF 1761
RESULT 7
CIN3 HUMAN STANDARD; PRT; 2000 AA.
ID CIN3_HUMAN Q9N46; Q16142; Q9B2B3; Q9C006; Q9MYK2; Q9UPD1; Q9YEP4;
AC Q9N46; Q16142; Q9B2B3; Q9C006; Q9MYK2; Q9UPD1; Q9YEP4;
DT 16-OCT-2001 (Rel. 40, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 01-OCT-2004 (Rel. 45, Last annotation update)
DE Sodium channel protein type III alpha subunit (Voltage-gated sodium channel alpha subunit Nav1.3) (Sodium channel protein, brain III alpha subunit) (Voltage-gated sodium channel subtype III).
DE subunit (Voltage-gated sodium channel subtype III).
GN Name=SCN3A; Synonyms=NA3;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A. (ISOFORM 2).
RC TISSUE=Brain;
RA Chen Y., Dale T.J., Romanos M.A., Whitaker W.R., Xie X., Clare J.J.;
RT "Cloning, distribution and functional analysis of the human brain type III sodium channel from human brain."
RL Submitted (DEC-1999) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A. (ISOFORM 3).
RA Jeong S.-Y., Goto J., Kanazawa I.;
RT "Cloning of cDNA for human voltage-gated sodium channel alpha subunit, SCN3A."
RL Submitted (JAN-2000) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A. (ISOFORMS 1; 2; 3 AND 4), AND VARIANT THR-606
RX MEDLINE=21142400; PubMed=11245585; DOI=10.1016/S0378-1119(00)00594-1;
RA Kasai N., Fukushima K., Ueki Y., Prasad S., Nosakowski J.,
RA Sugata K.-I., Sugata A., Nishizaki K., Meyer N.C., Smith R.J.H.;
RT "Genomic structures of SCN2A and SCN3A - candidate genes for deafness at the DFNA16 locus."
RL Gene 264:113-122(2001).
RN [4]
RP SEQUENCE OF 1-1415 FROM N.A. (ISOFORMS 2 AND 4).
RC TISSUE=Brain;
RX MEDLINE=98251277; PubMed=9589372;
RA Lu C.M., Brown G.B.;
RT "Isolation of a human-brain sodium-channel gene encoding two isoforms of the subtype III alpha-subunit."
RL J. Mol. Neurosci. 10:67-70(1998).
RN [5]
RP SEQUENCE OF 1324-1413 FROM N.A.
RC TISSUE=Placenta;
RX MEDLINE=94211784; PubMed=8159680;
RA Malo M.S., Striwasava K., Andreessen J.M., Chen X.N., Korenberg J.R., Ingram V.M.;
RT "Targeted gene walking by low stringency polymerase chain reaction: assignment of a putative human brain sodium channel gene (SCN3A) to chromosome 2q24-31."
RL Proc. Natl. Acad. Sci. U.S.A. 91:2875-2979(1994).
RN [6]
RP SEQUENCE OF 1669-1750 FROM N.A.
RC TISSUE=Kidney;
RA Tomkovich G.S., Kyle J.W.;
RT "Endogenous sodium current in HEK293 cells: increase in cell surface expression of endogenous currents by stable transfection of the Beta 1 subunit."
RL Submitted (FEB-2000) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: Mediates the voltage-dependent sodium ion permeability of excitable membranes. Assuming opened or closed conformations in

CC response to the voltage difference across the membrane, the
CC protein forms a sodium-selective channel through which Na(+) ions
CC may pass in accordance with their electrochemical gradient.
CC -1- SUBUNIT: The sodium channel consists of a large polypeptide and 2-
CC 3 smaller ones. This sequence represents a large polypeptide.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
CC -1- ALTERNATIVE PRODUCTS:
CC Event=Alternative splicing; Named isoforms=4;
CC Comment=Exons 6A and 6N only differ by a single residue;
CC Name=1; Synonyms=6A-12+12b;
CC IsoId=Q9NY46-1; Sequence=Displayed;
CC Name=2; Synonyms=6A-12;
CC IsoId=Q9NY46-2; Sequence=VSP_001034;
CC Name=3; Synonyms=6N-12+12b;
CC IsoId=Q9NY46-3; Sequence=VSP_001033;
CC Name=4; Synonyms=6N-12;
CC IsoId=Q9NY46-4; Sequence=VSP_001033, VSP_001034;
CC -1- DOMAIN: The sequence contains 4 internal repeats, each with 5
CC hydrophobic segments (S1,S2,S3,S5,S6) and one positively charged
CC segment (S4). Segments S4 are probably the voltage-sensors and are
CC characterized by a series of positively charged amino acids at
CC every third position.
CC -1- SIMILARITY: Belongs to the sodium channel family.
CC -1- SIMILARITY: Contains 1 IQ domain.
CC
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CC or send an email to license@sib-sib.ch).
CC
CC EMBL; AF251507; CAB85895.1; -;
CC EMBL; AF25987; AAK00219.1; -;
CC EMBL; AF330135; AAG53414.1; JOINED.
CC EMBL; AF330118; AAG53414.1; JOINED.
CC EMBL; AF330119; AAG53414.1; JOINED.
CC EMBL; AF330120; AAG53414.1; JOINED.
CC EMBL; AF330121; AAG53414.1; JOINED.
CC EMBL; AF330122; AAG53414.1; JOINED.
CC EMBL; AF330123; AAG53414.1; JOINED.
CC EMBL; AF330124; AAG53414.1; JOINED.
CC EMBL; AF330125; AAG53414.1; JOINED.
CC EMBL; AF330126; AAG53414.1; JOINED.
CC EMBL; AF330127; AAG53414.1; JOINED.
CC EMBL; AF330128; AAG53414.1; JOINED.
CC EMBL; AF330129; AAG53414.1; JOINED.
CC EMBL; AF330130; AAG53414.1; JOINED.
CC EMBL; AF330131; AAG53414.1; JOINED.
CC EMBL; AF330132; AAG53414.1; JOINED.
CC EMBL; AF330133; AAG53414.1; JOINED.
CC EMBL; AF330134; AAG53414.1; JOINED.
CC EMBL; AF330135; AAG53415.1; -;
CC EMBL; AF330118; AAG53415.1; JOINED.
CC EMBL; AF330119; AAG53415.1; JOINED.
CC EMBL; AF330120; AAG53415.1; JOINED.
CC EMBL; AF330121; AAG53415.1; JOINED.
CC EMBL; AF330122; AAG53415.1; JOINED.
CC EMBL; AF330123; AAG53415.1; JOINED.
CC EMBL; AF330124; AAG53415.1; JOINED.
CC EMBL; AF330125; AAG53415.1; JOINED.
CC EMBL; AF330126; AAG53415.1; JOINED.
CC EMBL; AF330127; AAG53415.1; JOINED.
CC EMBL; AF330128; AAG53415.1; JOINED.
CC EMBL; AF330129; AAG53415.1; JOINED.
CC EMBL; AF330130; AAG53415.1; JOINED.
CC EMBL; AF330131; AAG53415.1; JOINED.
CC EMBL; AF330132; AAG53415.1; JOINED.
CC EMBL; AF330133; AAG53415.1; JOINED.
CC EMBL; AF330134; AAG53415.1; JOINED.
CC EMBL; AF035685; AAC29514.1; -;
CC EMBL; AF035686; AAC29515.1; -;

DR EMBL; S69887; AAB30530.1; -
 DR EMBL; AF239921; AAP44690.1; -
 DR PIR; A54937; A54937.
 DR HSSP; P04775; 1BXY.
 DR Genew; HGNC:10590; SCN3A.
 DR MIM; 162391; -
 DR GO; GO:0001518; C:voltage-gated sodium channel complex; NAS.
 DR GO; GO:0005248; F:voltage-gated sodium channel activity; NAS.
 DR GO; GO:0006814; P:sodium ion transport; NAS.
 DR InterPro; IPR001682; Ca/Na pore.
 DR InterPro; IPR002111; Cat_channel_TripL.
 DR InterPro; IPR005821; Ion_trans.
 DR InterPro; IPR000048; Ion_region.
 DR InterPro; IPR005820; M+channel_nlg.
 DR InterPro; IPR001696; Na_channel.
 DR Pfam; PF00520; Ion_trans; 4.
 DR Pfam; PF06512; IQ_1.
 DR Pfam; PF06512; Na_trans_assoc; 1.
 DR PRINTS; PR00170; NACHANNEL.
 DR PROSITE; PS50096; IQ_1.
 KW Alternative splicing; Glycoprotein; Ion transport; Ionic channel;
 KW Multigene family; Polymorphism; Repeat; Sodium channel; Transmembrane;
 KW Voltage-gated channel.
 FT TRANSMEM 124 147 S1 of repeat 1.
 FT TRANSMEM 156 175 S2 of repeat 1.
 FT TRANSMEM 189 207 S3 of repeat 1.
 FT TRANSMEM 214 233 S4 of repeat 1.
 FT TRANSMEM 249 273 S5 of repeat 1.
 FT TRANSMEM 401 426 S6 of repeat 1.
 FT TRANSMEM 455 479 S1 of repeat 1.
 FT TRANSMEM 755 779 S2 of repeat 1.
 FT TRANSMEM 791 814 S3 of repeat 1.
 FT TRANSMEM 823 842 S4 of repeat 1.
 FT TRANSMEM 849 869 S5 of repeat 1.
 FT TRANSMEM 885 905 S6 of repeat 1.
 FT TRANSMEM 959 984 S1 of repeat 1.
 FT TRANSMEM 1202 1225 S2 of repeat 1.
 FT TRANSMEM 1239 1264 S3 of repeat 1.
 FT TRANSMEM 1271 1292 S4 of repeat 1.
 FT TRANSMEM 1318 1338 S5 of repeat 1.
 FT TRANSMEM 1338 1359 S6 of repeat 1.
 FT TRANSMEM 1442 1468 S1 of repeat 1.
 FT TRANSMEM 1522 1545 S2 of repeat 1.
 FT TRANSMEM 1557 1580 S3 of repeat 1.
 FT TRANSMEM 1587 1610 S4 of repeat 1.
 FT TRANSMEM 1621 1642 S5 of repeat 1.
 FT TRANSMEM 1658 1680 S6 of repeat 1.
 FT TRANSMEM 1747 1771 S1 of repeat 1.
 FT TRANSMEM 1900 1929 S2 of repeat 1.
 FT TRANSMEM 211 231 S3 of repeat 1.
 FT TRANSMEM 290 310 S4 of repeat 1.
 FT TRANSMEM 296 316 S5 of repeat 1.
 FT TRANSMEM 302 322 S6 of repeat 1.
 FT TRANSMEM 307 327 S1 of repeat 1.
 FT TRANSMEM 339 359 S2 of repeat 1.
 FT TRANSMEM 364 384 S3 of repeat 1.
 FT TRANSMEM 424 444 S4 of repeat 1.
 FT TRANSMEM 484 504 S5 of repeat 1.
 FT TRANSMEM 1051 1071 S6 of repeat 1.
 FT TRANSMEM 1068 1088 S1 of repeat 1.
 FT TRANSMEM 1134 1154 S2 of repeat 1.
 FT TRANSMEM 1366 1386 S3 of repeat 1.
 FT TRANSMEM 1380 1400 S4 of repeat 1.
 FT TRANSMEM 208 228 S5 of repeat 1.
 FT TRANSMEM 673 693 S6 of repeat 1.
 FT VARSPLIC 673 693 S1 of repeat 1.
 Query Match 100.0%; Score 28; DB 1; Length 2000;
 Best Local Similarity 100.0%; Pred. No. 67e-19;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GIFFEVSYIIISFLVVMYTAIVLENP 28
 DB 1747 GIFFEVSYIIISFLVVMYTAIVLENP 1774

RESULT 8
 ID CIN2_HUMAN STANDARD; PRT; 2005 AA.
 AC Q99250; Q99250; Q99250; Q99250;
 DT 01-JUN-1994 (Rel. 29, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 05-JUL-2004 (Rel. 44, Last annotation update)
 DE channel protein type II alpha subunit (Voltage-gated sodium
 DE channel alpha subunit Nav1.2) (Sodium channel protein, brain II alpha
 DE subunit) (HSC II)
 GN Homo sapiens (Human)
 OS Homo sapiens (Human)
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
 OX NCBI_Taxid=9606;
 RN [1]
 RP SEQUENCE FROM N.A. (ISOFORM 1).
 RC TISSUE=Brain;
 RX MEDLINE=92390418; PubMed=1325650;
 RA Ahmed C.M., Ware D.H., Lee S.C., Patten C.D., Ferrer-Montiel A.V.,
 RA Schinder A.F., McPherson J.D., Wagner-McPherson C.B., Wasmuth J.J.,
 RA Evans G.A., Montiel M.;
 RT "Primary structure, chromosomal localization, and functional
 RT expression of a voltage-gated sodium channel from human brain.";
 RT Proc. Natl. Acad. Sci. U.S.A. 89:8220-8224(1992).
 RN [2]
 RP SEQUENCE FROM N.A. (ISOFORMS 1 AND 2).
 RC MEDLINE=21142400; PubMed=11245985; DOI=10.1016/S0378-1119(00)00594-1;
 RA Kasai N., Fukushima K., Ueki Y., Prasad S., Nosakowski J.,
 RA Sugata K.-I., Sugata A., Nishizaki K., Meyer N.C., Smith R.J.H.,
 RA "Genomic structures of SCN2A and SCN3A - candidate genes for deafness
 RA at the DFNA16 locus.";
 RT Gene 264:113-122(2001).
 RN [3]
 RP SEQUENCE OF 1-89 FROM N.A.
 RC Lu C.-M., Eichelsberger U.S., Beckman M.L., Schade S.D., Brown G.B.;
 RA "Isolation of the 5'-flanking region for human brain sodium channel
 RA subtype II alpha-subunit (SCN2A).";
 RT Submitted (Apr-1998) to the EMBL/Genbank/DBJ databases.
 RN [4]
 RP SEQUENCE OF 1702-2005 FROM N.A.
 RC TISSUE=Brain;
 RX MEDLINE=92275082; PubMed=1317301;
 RA Lu C.-M., Han U., Rado T.A., Brown G.B.;
 RA "Differential expression of two sodium channel subtypes in human
 RA brain.";
 RT FEBS Lett. 303:53-58(1992).
 RN [5]
 RP SEQUENCE OF 1702-1772 FROM N.A.
 RX MEDLINE=91110524; PubMed=1846440;
 RA Han U., Lu C.-M., Brown G.B., Rado T.A.;
 RT "Direct amplification of a single dissected chromosomal segment by
 RT polymerase chain reaction: a human brain sodium channel gene is on
 RT chromosome 2q22-q23.";
 RT Proc. Natl. Acad. Sci. U.S.A. 88:335-339(1991).
 RL -1- FUNCTION: Mediates the voltage-dependent sodium ion permeability
 RL of excitable membranes. Assuming opened or closed conformations in
 RL response to the voltage difference across the membrane, the
 RL protein forms a sodium-selective channel through which Na(+) ions
 RL may pass in accordance with their electrochemical gradient.
 RL -1- SUBUNIT: The sodium channel consists of a large polypeptide and 2-
 RL 3 smaller ones. This sequence represents a large polypeptide.
 RL -1- SUBCELLULAR LOCATION: Integral membrane protein.
 RL -1- ALTERNATIVE PRODUCTS:
 RL Event=Alternative splicing; Named isoforms=2;
 RL Name=1; Synonyms=Adult, 6A;
 RL Name=2; Synonyms=Neonatal, 6N;
 RL IsoId=Q99250-1; Sequence=Displayed;
 RL IsoId=Q99250-2; Sequence=VSP_001032;
 RL -1- DOMAIN: The sequence contains 4 internal repeats, each with 5
 RL hydrophobic segments (S1,S2,S3,S5,S6) and one positively charged

segment (S4). Segments S4 are probably the voltage-sensors and are characterized by a series of positively charged amino acids at every third position.

-1- SIMILARITY: Belongs to the sodium channel family.

-1- SIMILARITY: Contains 1 IQ domain.

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CC -----

DR EMBL, M94055; AAA18895.1; -.

DR EMBL, AF059683; AAC14574.1; -.

DR EMBL, AF327246; AAG53413.1; JOINED.

DR EMBL, AF327227; AAG53413.1; JOINED.

DR EMBL, AF327228; AAG53413.1; JOINED.

DR EMBL, AF327229; AAG53413.1; JOINED.

DR EMBL, AF327230; AAG53413.1; JOINED.

DR EMBL, AF327231; AAG53413.1; JOINED.

DR EMBL, AF327232; AAG53413.1; JOINED.

DR EMBL, AF327233; AAG53413.1; JOINED.

DR EMBL, AF327234; AAG53413.1; JOINED.

DR EMBL, AF327235; AAG53413.1; JOINED.

DR EMBL, AF327236; AAG53413.1; JOINED.

DR EMBL, AF327237; AAG53413.1; JOINED.

DR EMBL, AF327238; AAG53413.1; JOINED.

DR EMBL, AF327239; AAG53413.1; JOINED.

DR EMBL, AF327240; AAG53413.1; JOINED.

DR EMBL, AF327241; AAG53413.1; JOINED.

DR EMBL, AF327242; AAG53413.1; JOINED.

DR EMBL, AF327243; AAG53413.1; JOINED.

DR EMBL, AF327244; AAG53413.1; JOINED.

DR EMBL, AF327245; AAG53413.1; JOINED.

DR EMBL, AF327246; AAG53412.1; -.

DR EMBL, AF327226; AAG53412.1; JOINED.

DR EMBL, AF327227; AAG53412.1; JOINED.

DR EMBL, AF327228; AAG53412.1; JOINED.

DR EMBL, AF327229; AAG53412.1; JOINED.

DR EMBL, AF327230; AAG53412.1; JOINED.

DR EMBL, AF327231; AAG53412.1; JOINED.

DR EMBL, AF327232; AAG53412.1; JOINED.

DR EMBL, AF327233; AAG53412.1; JOINED.

DR EMBL, AF327234; AAG53412.1; JOINED.

DR EMBL, AF327235; AAG53412.1; JOINED.

DR EMBL, AF327236; AAG53412.1; JOINED.

DR EMBL, AF327237; AAG53412.1; JOINED.

DR EMBL, AF327238; AAG53412.1; JOINED.

DR EMBL, AF327239; AAG53412.1; JOINED.

DR EMBL, AF327240; AAG53412.1; JOINED.

DR EMBL, AF327241; AAG53412.1; JOINED.

DR EMBL, AF327242; AAG53412.1; JOINED.

DR EMBL, AF327243; AAG53412.1; JOINED.

DR EMBL, AF327244; AAG53412.1; JOINED.

DR EMBL, AF327245; AAG53412.1; JOINED.

DR EMBL, M91804; -; NOT ANNOTATED_CDS.

DR EMBL, M55662; AAB65854.2; -.

DR HSSP, P04775; 1BYX; SCN2A2.

DR Genew; HGNC:10588; SCN2A2.

DR MIM; 601219; -.

DR GO; GO:0005887; C:integral to plasma membrane; TAS.

DR GO; GO:0005248; P:voltage-gated sodium channel activity; TAS.

DR GO; GO:0006814; P:sodium ion transport; TAS.

DR InterPro; IPR001682; Ca/Na_pore.

DR InterPro; IPR002111; Cat_channel_TrpL.

DR InterPro; IPR005821; Ion_trans.

DR InterPro; IPR000048; IQ_region.

DR InterPro; IPR005820; M+channel_nlg.

DR InterPro; IPR001696; Na_channel.

DR InterPro; IPR010526; Na_trans_assoc.

DR Pfam; PF005020; Ion_trans; 4.

DR Pfam; PF06512; IQ; 1.

DR Pfam; PF06512; Na_trans_assoc; 1.

DR PRINTS; PR00170; NACHANNEL.

DR PROSITE; PS50096; IQ; 1.

KW Alternative splicing; Glycoprotein; Ion transport; Ionic channel;

KW Multigene family; Repeat; Sodium channel; Transmembrane;

KW Voltage-gated channel.

FT REPEAT 111 456 I.

FT REPEAT 741 1013 II.

FT REPEAT 1190 1504 III.

FT REPEAT 1513 1811 IV.

FT TRANSMEM 125 148 S1 of repeat I.

FT TRANSMEM 157 176 S2 of repeat I.

FT TRANSMEM 190 208 S3 of repeat I.

FT TRANSMEM 215 234 S4 of repeat I.

FT TRANSMEM 251 274 S5 of repeat I.

FT TRANSMEM 402 427 S6 of repeat I.

FT TRANSMEM 754 778 S1 of repeat II.

FT TRANSMEM 790 813 S2 of repeat II.

FT TRANSMEM 822 841 S3 of repeat II.

FT TRANSMEM 848 867 S4 of repeat II.

FT TRANSMEM 884 904 S5 of repeat II.

FT TRANSMEM 958 983 S6 of repeat II.

FT TRANSMEM 1204 1227 S1 of repeat III.

FT TRANSMEM 1241 1266 S2 of repeat III.

FT TRANSMEM 1273 1294 S3 of repeat III.

FT TRANSMEM 1299 1320 S4 of repeat III.

FT TRANSMEM 1340 1367 S5 of repeat III.

FT TRANSMEM 1447 1473 S6 of repeat III.

FT TRANSMEM 1527 1550 S1 of repeat IV.

FT TRANSMEM 1592 1585 S2 of repeat IV.

FT TRANSMEM 1592 1615 S3 of repeat IV.

FT TRANSMEM 1626 1647 S4 of repeat IV.

FT TRANSMEM 1663 1685 S5 of repeat IV.

FT TRANSMEM 1752 1766 S6 of repeat IV.

FT DOMAIN 1905 1934 IQ.

FT CARBOHYD 212 212 N-linked (GlcNAc . .) (Potential).

FT CARBOHYD 285 285 N-linked (GlcNAc . .) (Potential).

FT CARBOHYD 291 291 N-linked (GlcNAc . .) (Potential).

FT CARBOHYD 297 297 N-linked (GlcNAc . .) (Potential).

FT CARBOHYD 303 303 N-linked (GlcNAc . .) (Potential).

FT CARBOHYD 308 308 N-linked (GlcNAc . .) (Potential).

FT CARBOHYD 340 340 N-linked (GlcNAc . .) (Potential).

FT CARBOHYD 604 604 N-linked (GlcNAc . .) (Potential).

FT CARBOHYD 624 624 N-linked (GlcNAc . .) (Potential).

FT CARBOHYD 883 883 N-linked (GlcNAc . .) (Potential).

FT CARBOHYD 1055 1055 N-linked (GlcNAc . .) (Potential).

FT CARBOHYD 1072 1072 N-linked (GlcNAc . .) (Potential).

FT CARBOHYD 1136 1136 N-linked (GlcNAc . .) (Potential).

FT CARBOHYD 1368 1368 N-linked (GlcNAc . .) (Potential).

FT CARBOHYD 1382 1382 N-linked (GlcNAc . .) (Potential).

FT CARBOHYD 1393 1393 N-linked (GlcNAc . .) (Potential).

FT CARBOHYD 1778 1778 N-linked (GlcNAc . .) (Potential).

FT VARSPLIC 209 209 D -> N (1n isoform 2).

FT CONFLICT 524 524 /FTId=VSP_001032.

R -> L (1n Ref. 1).

Query Match 100.0%; Score 28; DB 1; Length 2005;

Best Local Similarity 100.0%; Pred. No. 6.7e-19;

Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GIFFFVSYIIIFLVVNNYIVIIENF 28

Db 1752 GIFFFVSYIIIFLVVNNYIVIIENF 1779

RESULT 9

ID CIN2 RAT STANDARD; PRT; 2005 AA.

AC P04775;

DT 13-AUG-1987 (Rel. 05, Created)

DT 13-AUG-1987 (Rel. 05, Last sequence update)
 DT 01-OCT-2004 (Rel. 45, Last annotation update)
 DE Sodium channel protein type II alpha subunit (Voltage-gated sodium
 DE channel alpha subunit Nav1.2) (Sodium channel protein, brain II alpha
 DE subunit).
 GN Name=Scn2a;
 OS Rattus norvegicus (Rat).
 CC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 CC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 ON NCBI_TaxID=10116;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA MEDLINE=86146901; PubMed=3754035;
 RA Noda M., Ikeda T., Kayano T., Suzuki H., Takeshima H., Kuraaki M.,
 RA Takahashi H., Numa S.;
 RL "Existence of distinct sodium channel messenger RNAs in rat brain.",
 CC Nature 320:186-192(1986).
 CC -1- FUNCTION: Mediates the voltage-dependent sodium ion permeability
 CC of excitable membranes. Assuming opened or closed conformations in
 CC response to the voltage difference across the membrane, the
 CC protein forms a sodium-selective channel through which Na(+) ions
 CC may pass in accordance with their electrochemical gradient.
 CC -1- SUBUNIT: The sodium channel consists of a large polypeptide and 2-
 CC 3 smaller ones. This sequence represents a large polypeptide.
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
 CC hydrophobic segments (S1, S2, S3, S5, S6) and one positively charged
 CC segment (S4). Segments S4 are probably the voltage-sensors and are
 CC characterized by a series of positively charged amino acids at
 CC every third position.
 CC -1- SIMILARITY: Belongs to the sodium channel family.
 CC -1- SIMILARITY: Contains 1 IQ domain.
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 CC or send an email to license@sib-sib.ch).
 CC -----
 DR EMBL: X03639; CAA27287.1; -
 DR PDB: 1BY7; NMR: A=1474-1526.
 DR InterPro: IPR001682; Ca/Na_Pore.
 DR InterPro: IPR002111; Cat_channel_TripL.
 DR InterPro: IPR005821; Ion_trans.
 DR InterPro: IPR000048; IQ_region.
 DR InterPro: IPR005820; M-channel_nlg.
 DR InterPro: IPR01656; Na_channel.
 DR Pfam: PF00520; Ion_trans; 4.
 DR Pfam: PF00612; IQ; 1.
 DR Pfam: PF06512; Na_trans_assoc; 1.
 DR PRINTS: PR00170; NACHANNEL.
 DR PROSITE: PS50096; IQ; 1
 DR 3D-structure; Glycoprotein; Ion transport; Ionic channel;
 KM Multigene family; Repeat; Sodium channel; Transmembrane;
 KM Voltage-gated channel.
 FT REPEAT 111 456 I.
 FT REPEAT 741 1013 II.
 FT REPEAT 1190 1504 III.
 FT REPEAT 1513 1811 IV.
 FT TRANSMEM 125 148 S1 of repeat I.
 FT TRANSMEM 157 176 S2 of repeat I.
 FT TRANSMEM 190 208 S3 of repeat I.
 FT TRANSMEM 215 234 S4 of repeat I.
 FT TRANSMEM 251 274 S5 of repeat I.
 FT TRANSMEM 402 427 S6 of repeat I.
 FT TRANSMEM 754 778 S1 of repeat II.
 FT TRANSMEM 790 813 S2 of repeat II.
 FT TRANSMEM 822 841 S3 of repeat II.
 FT TRANSMEM 848 867 S4 of repeat II.
 FT TRANSMEM 884 904 S5 of repeat II.

FT TRANSMEM 958 983 S6 of repeat II.
 FT TRANSMEM 1204 1227 S1 of repeat III.
 FT TRANSMEM 1241 1266 S2 of repeat III.
 FT TRANSMEM 1273 1294 S3 of repeat III.
 FT TRANSMEM 1299 1320 S4 of repeat III.
 FT TRANSMEM 1340 1367 S5 of repeat III.
 FT TRANSMEM 1447 1473 S6 of repeat III.
 FT TRANSMEM 1527 1550 S1 of repeat IV.
 FT TRANSMEM 1562 1585 S2 of repeat IV.
 FT TRANSMEM 1592 1615 S3 of repeat IV.
 FT TRANSMEM 1626 1647 S4 of repeat IV.
 FT TRANSMEM 1663 1685 S5 of repeat IV.
 FT TRANSMEM 1752 1776 S6 of repeat IV.
 FT DOMAIN 1905 1934 IQ.
 FT CARBOHYD 212 212 N-linked (GlcNAc...)
 FT CARBOHYD 285 285 N-linked (GlcNAc...)
 FT CARBOHYD 291 291 N-linked (GlcNAc...)
 FT CARBOHYD 297 297 N-linked (GlcNAc...)
 FT CARBOHYD 303 303 N-linked (GlcNAc...)
 FT CARBOHYD 308 308 N-linked (GlcNAc...)
 FT CARBOHYD 340 340 N-linked (GlcNAc...)
 FT CARBOHYD 604 604 N-linked (GlcNAc...)
 FT CARBOHYD 624 624 N-linked (GlcNAc...)
 FT CARBOHYD 883 883 N-linked (GlcNAc...)
 FT CARBOHYD 1055 1055 N-linked (GlcNAc...)
 FT CARBOHYD 1072 1072 N-linked (GlcNAc...)
 FT CARBOHYD 1136 1136 N-linked (GlcNAc...)
 FT CARBOHYD 1168 1168 N-linked (GlcNAc...)
 FT CARBOHYD 1382 1382 N-linked (GlcNAc...)
 FT CARBOHYD 1393 1393 N-linked (GlcNAc...)
 SQ SEQUENCE 2005 AA; 227872 MW; 861B583D79F8324 CRC64;
 Query Match 100.0%; Score 28; DB 1; Length 2005;
 Best Local Similarity 100.0%; Pred. No. 6,7e-19;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 GIFFPVSIIISFLVVMYAVILENF 28
 Db 1752 GIFFPVSIIISFLVVMYAVILENF 1779
 RESULT 10
 Q9YGN7 PRELIMINARY; PRT; 2007 AA.
 AC Q9YGN7;
 DT 01-MAY-1999 (TrEMBLrel. 10, Created)
 DT 01-MAY-1999 (TrEMBLrel. 10, Last sequence update)
 DE 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
 DE Voltage-dependent sodium channel.
 OS Cynops pyrrhogaster (Japanese common newt).
 CC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 CC Amphibia; Batrachia; Caudata; Salamandridae; Cynops.
 ON NCBI_TaxID=8330;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA TISUB-Retina;
 RA Hirota K., Kaneko Y., Matsumoto G., Hanyu Y.;
 RL Submitted (JAN-1999) to the EMBL/GenBank/DBJ databases.
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
 DR EMBL: AF123593; AADI7315.1; -.
 DR HSRF: P04775; 1BY7.
 DR GO: GO:0016021; C:Integral to membrane; IEA.
 DR GO: GO:001518; C:voltage-gated sodium channel complex; IEA.
 DR GO: GO:0005261; F:cation channel activity; IEA.
 DR GO: GO:0005248; F:voltage-gated sodium channel activity; IEA.
 DR GO: GO:0006812; P:cation transport; IEA.
 DR GO: GO:0006814; P:sodium ion transport; IEA.
 DR InterPro: IPR001682; Ca/Na_Pore.
 DR InterPro: IPR002111; Cat_channel_TripL.
 DR InterPro: IPR005821; Ion_trans.
 DR InterPro: IPR005820; M-channel_nlg.

DR InterPro; IPRO01696; Na_channel.
DR InterPro; IPRO10526; Na_trans_assoc.
DR Pfam; PF00612; Ion_trans_4.
DR Pfam; PF00612; IQ_1.
DR Pfam; PF06512; Na_trans_assoc; 1.
DR PRINTS; PRO0170; NACHANNEL.
DR SMART; SM00015; IQ_1.
KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
KW Transport; Voltage-gated channel.
SQ SEQUENCE 2007 AA; 228398 MW; 013B9B9EC9C94C9 CRC64;

Query Match 100.0%; Score 28; DB 2; Length 2007;
Best Local Similarity 100.0%; Pred. NO. 6.7e-19;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GIFFVSYIIISFLVVMNYIAVILENF 28
DB 1754 GIFFVSYIIISFLVVMNYIAVILENF 1781

RESULT 11
CINI_HUMAN STANDARD; PRT; 2009 AA.
ID CINI_HUMAN
AC P35498; Q16172; Q96LA3; Q9C008;
DT 01-JUN-1994 (Rel. 29, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 05-JUL-2004 (Rel. 44, Last annotation update)
DE Sodium channel protein type I alpha subunit (Voltage-gated sodium
DE channel alpha subunit Nav1.1) (Sodium channel protein, brain I alpha
DE subunit).
GN Name=SCN1A; Synonym=SCN1, NAC1;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_TaxID=9606;
SQ SEQUENCE FROM N.A. (ISOFORM 1), AND VARIANTS GEFS+2 MET-875 AND
RP HIS-1648.
RX MEDLINE=20206553; PubMed=10742094;
RA Becay A., MacDonald B.T., Meisler M.H., Baulac S., Huberfeld G.,
RA An-Goufinkel I., Brice A., LeGuern E., Mouldard B., Chaigne D.,
RA Buresi C., Malafosse A.;
RT "Mutations of SCN1A, encoding a neuronal sodium channel, in two
RT families with GEFS+2.";
RL Nat. Genet. 24:343-345(2000).
RN
RN SEQUENCE FROM N.A. (ISOFORM 2).
RA Jeong S.-Y., Goto J., Kanazawa I.;
RT "Cloning of cDNA for human voltage-gated sodium channel alpha subunit,
RT SCN1A.";
RL Submitted (JAN-2000) to the EMBL/GenBank/DBJ databases.
RN
RN SEQUENCE FROM N.A. (ISOFORM 2).
RC TISSUE=Brain;
RA Sugawara T., Mazaki E.M., Yamakawa K.;
RT "Homo sapiens neuronal voltage-gated sodium channel type I (Nav1.1)
RT mRNA.";
RL Submitted (JUL-2001) to the EMBL/GenBank/DBJ databases.
RN
RN SEQUENCE FROM N.A. (ISOFORMS 1 AND 2), AND VARIANT THR-1067.
RP Ouchida M., Omori I.;
RT "Isoforms of human sodium channel SCN1A gene.";
RL Submitted (OCT-2002) to the EMBL/GenBank/DBJ databases.
RN
RN SEQUENCE OF 1335-1428 FROM N.A.
RX MEDLINE=94340991; PubMed=8062593;
RA Malo M.S., Blanchard B.J., Andresen J.M., Srivastava K., Chen X.N.,
RA Li X., Jabe E.W., Korenberg J.R., Ingram V.M.;
RT "Localization of a putative human brain sodium channel gene (SCN1A) to
RT chromosome band 2q24.";
RL Cytogenet. Cell Genet. 67:178-186(1994).
RN
RN SEQUENCE OF 1518-1940 FROM N.A.

RC TISSUE=Brain;
RX MEDLINE=92275082; PubMed=1317301;
RA Lu C.-M., Han J., Rado T.A., Brown G.B.;
RT "Differential expression of two sodium channel subtypes in human
RT brain.";
RL FEBS Lett. 303:53-58(1992).
RN
RN VARIANTS GEFS+2 VAL-188; LEU-1353 AND MET-1656.
RX MEDLINE=21152275; PubMed=11254444;
RA Wallace R.H., Scheffer I.E., Barnett S., Richards M., Dibbens L.,
RA Desai R.R., Lerman-Sagie T., Lev D., Mazarib A., Brand N.,
RA Ben-Zeev B., Golikman I., Singh R., Kremmidiotis G., Gardner A.,
RA Sutherland G.R., George A.L. Jr., Mulley J.C., Berkovic S.F.;
RT "Neuronal sodium-channel alpha1-subunit mutations in generalized
RT epilepsy with febrile seizures plus.";
RL Am. J. Hum. Genet. 68:859-865(2001).
RN
RN VARIANT GEFS+2 ARG-1204.
RX MEDLINE=21152275; PubMed=11254445;
RA Becay A., Heils A., MacDonald B.T., Haug K., Sander T., Meisler M.H.;
RT "A novel SCN1A mutation associated with generalized epilepsy with
RT febrile seizures plus -- and prevalence of variants in patients with
RT epilepsy.";
RL Am. J. Hum. Genet. 68:866-873(2001).
RN
RN VARIANT SMEI PHE-986.
RX MEDLINE=21257503; PubMed=11359211;
RA Claes L., Del-Favero J., Ceulemans B., Lagae L., Van Broeckhoven C.,
RA De Jonghe P.;
RT "De novo mutations in the sodium-channel gene SCN1A cause severe
RT myoclonic epilepsy of infancy.";
RL Am. J. Hum. Genet. 68:1327-1332(2001).
RN
RN VARIANT GEFS+2 THR-1270.
RX MEDLINE=21630138; PubMed=11756608;
RA Abou-Khalil B., Ge Q., Desai R., Ryther R., Bazyk A., Bailey R.,
RA Haines J.L., Sutcliffe J.S., George A.L. Jr.;
RT "Partial and generalized epilepsy with febrile seizures plus and a
RT novel SCN1A mutation.";
RL Neurology 57:2265-2272(2001).
RN
RN FUNCTION: Mediates the voltage-dependent sodium ion permeability
RN of excitable membranes. Assuming opened or closed conformations in
RN response to the voltage difference across the membrane, the
RN protein forms a sodium-selective channel through which Na(+) ions
RN may pass in accordance with their electrochemical gradient.
RN SUBUNIT: The sodium channel consists of a large polypeptide and 2-
RN 3 smaller ones. This sequence represents a large polypeptide.
RN -1- SUBCELLULAR LOCATION: Integral membrane protein.
RN -1- ALTERNATIVE PRODUCTS:
RN Event=Alternative splicing; Named isoforms=2;
RN Name=1;
RN IsoId=P35498-1; Sequence=Displayed;
RN Name=2;
RN IsoId=P35498-2; Sequence=VSP_001031;
RN Note=No experimental confirmation available;
RN -1- DOMAIN: The sequence contains 4 internal repeats, each with 5
RN hydrophobic segments (S1,S2,S3,S5,S6) and one positively charged
RN segment (S4). Segments S4 are probably the voltage-sensors and are
RN characterized by a series of positively charged amino acids at
RN every third position.
RN -1- DISEASE: Defects in SCN1A are the cause of generalized epilepsy
RN with febrile seizures plus type 2 (GEFS+2) [MIM:604233]. This
RN autosomal dominant disorder is characterized by febrile seizures
RN in children and afebrile seizures in adults. Penetrance is
RN incomplete and a large intrafamilial variability of the phenotype
RN is observed.
RN -1- DISEASE: Defects in SCN1A are a cause of severe myoclonic epilepsy
RN in infancy (SMEI) [MIM:607208], a severe form of generalized
RN epilepsy with febrile seizures. SMEI is a rare disorder
RN characterized by normal development before onset, seizures
RN beginning in the first year of life in the form of generalized or
RN unilateral febrile clonic seizures, secondary appearance of
RN myoclonic seizures, and occasionally partial seizures. It is

CC associated with ataxia, slowed psychomotor development, and mental decline.
 CC -1- SIMILARITY: Belongs to the sodium channel family.
 CC -1- SIMILARITY: Contains 1 IQ domain.
 CC -----
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 CC or send an email to license@isb-sib.ch).
 CC -----
 DR EMBL; AF225985; AAK00217.1; -;
 DR EMBL; AY043484; AAK95360.1; -;
 DR EMBL; AB093548; BAC21101.1; -;
 DR EMBL; AB093549; BAC21102.1; -;
 DR EMBL; S71446; AAB31605.1; -;
 DR EMBL; X65362; CAA46439.1; -;
 DR EMBL; M91803; -; NOT_ANNOTATED_CDS.
 DR PIR; I52964; I52964.
 DR PIR; S29184; S29184.
 DR HSSP; P04775; IBY.
 DR Gene; HGNC:10585; SCN1A.
 DR MIM; 182389; -;
 DR MIM; 604203; -;
 DR MIM; 607208; -;
 DR GO; GO:0016021; C:integral to membrane; NAS.
 DR GO; GO:0005248; F:voltage-gated sodium channel activity; NAS.
 DR GO; GO:0006814; P:sodium ion transport; NAS.
 DR InterPro; IPR001682; Ca/Na_pore.
 DR InterPro; IPR002111; Cat_channel_TpPL.
 DR InterPro; IPR005821; Ion_trans.
 DR InterPro; IPR000048; IQ_region.
 DR InterPro; IPR001696; M_channel_nlg.
 DR InterPro; IPR008051; Na_channel.
 DR InterPro; IPR010526; Na_trans_assoc.
 DR Pfam; PF00520; Ion_trans_4.
 DR Pfam; PF00612; IQ_1.
 DR Pfam; PF06512; Na_trans_assoc; 1.
 DR PRINTS; PR00170; NACHANNEL.
 DR PRINTS; PR01664; NACHANNEL.
 DR PROSITE; PS50096; IQ; FALSE_NEG.
 DR KW Alternative splicing; Disease mutation; Epilepsy; Glycoprotein;
 DR Ion transport; Ionic channel; Multigene family; Polymorphism; Repeat;
 DR Sodium channel; Transmembrane; Voltage-gated channel.
 FT REPEAT 110 454
 FT REPEAT 750 1022
 FT REPEAT 1200 1514
 FT REPEAT 1523 1821
 FT TRANSMEM 124 147
 FT TRANSMEM 156 175
 FT TRANSMEM 189 207
 FT TRANSMEM 214 223
 FT TRANSMEM 250 273
 FT TRANSMEM 400 425
 FT TRANSMEM 763 787
 FT TRANSMEM 799 822
 FT TRANSMEM 831 850
 FT TRANSMEM 857 876
 FT TRANSMEM 893 913
 FT TRANSMEM 967 992
 FT TRANSMEM 1214 1237
 FT TRANSMEM 1251 1276
 FT TRANSMEM 1283 1304
 FT TRANSMEM 1309 1330
 FT TRANSMEM 1350 1377
 FT TRANSMEM 1457 1483
 FT TRANSMEM 1537 1560
 FT TRANSMEM 1572 1595
 FT TRANSMEM 1602 1625
 FT TRANSMEM 1636 1657

FT TRANSMEM 1673 1695 S5 of repeat IV (By similarity).
 FT TRANSMEM 1762 1786 S6 of repeat IV (By similarity).
 FT CARBOHYD 211 211 N-linked (GlcNAc...) (potential).
 Query Match 100.0%; Score 28; DB 1; Length 2009;
 Best Local Similarity 100.0%; Pred. No. 6; 7e-19;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 GIFFVSYIIISFLVVMNYIAVILENF 28
 DB 1762 GIFFVSYIIISFLVVMNYIAVILENF 1789
 RESULT 12
 ID CINI_RAT STANDARD; PRT; 2009 AA.
 AC P04774;
 DT 13-AUG-1987 (Rel. 05, Created)
 DT 13-AUG-1987 (Rel. 05, Last sequence update)
 DT 05-JUL-2004 (Rel. 44, Last annotation update)
 DE Sodium channel protein type I alpha subunit (Voltage-gated sodium
 DE channel alpha subunit Nav1.1) (Sodium channel protein, brain I alpha
 DE subunit).
 DE GN Name=Scn1a;
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 OX NCBI_TaxId=10116;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA MEDLINE=86146901; PubMed=3754035;
 RA Noda M., Ikeda T., Kayano T., Suzuki H., Takeshima H., Kuraaki M.,
 RA Takahashi H., Numa S.;
 RT "Existence of distinct sodium channel messenger RNAs in rat brain.";
 RL Nature 320:188-192(1986).
 RN [2]
 RP SEQUENCE FROM N.A.
 RA MEDLINE=87313395; PubMed=2442385;
 RA Noda M., Numa S.;
 RT "Structure and function of sodium channel.";
 RL J. Recept. Res. 7:467-497(1987).
 CC -1- FUNCTION: Mediates the voltage-dependent sodium ion permeability
 CC of excitable membranes. Assuming opened or closed conformations in
 CC response to the voltage difference across the membrane, the
 CC protein forms a sodium-selective channel through which Na(+) ions
 CC may pass in accordance with their electrochemical gradient.
 CC -1- SUBUNIT: The sodium channel consists of a large polypeptide and 2-
 CC 3 smaller ones. This sequence represents a large polypeptide.
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
 CC -1- DOMAIN: The sequence contains 4 internal repeats, each with 5
 CC hydrophobic segments (S1, S2, S3, S5, S6) and one positively charged
 CC segment (S4). Segments S4 are probably the voltage-sensors and are
 CC characterized by a series of positively charged amino acids at
 CC every third position.
 CC -1- SIMILARITY: Belongs to the sodium channel family.
 CC -1- SIMILARITY: Contains 1 IQ domain.
 CC -----
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 CC -----
 DR EMBL; X03638; CAA27286.1; -;
 DR EMBL; M22253; AAK79965.1; -;
 DR PIR; A25019; A25019.
 DR HSSP; P04775; IBY.
 DR RGD; 69364; Scn1a.
 DR InterPro; IPR001682; Ca/Na_pore.
 DR InterPro; IPR002111; Cat_channel_TpPL.
 DR InterPro; IPR005821; Ion_trans.

DR InterPro; IPR000048; IQ region.
 DR InterPro; IPR005820; M+channel nlg.
 DR InterPro; IPR001696; Na channel.
 DR InterPro; IPR008051; Na channel.
 DR InterPro; IPR010526; Na_trans_assoc.
 DR Pfam; PF00520; Ion_trans_4.
 DR Pfam; PF06512; IQ_1.
 DR Pfam; PF06512; Na_trans_assoc; 1.
 DR PRINTS; PRO0170; NACHANNEL.
 DR PRINTS; PRO1664; NACHANNEL.
 DR PROSITE; PS50096; IQ; FALSE_NEG.
 KW Glycoprotein; Ion transport; Ionic channel; Multigene family; Repeat;
 KW Sodium channel; Transmembrane; Voltage-gated channel.
 FT REPEAT 110 454 I.
 FT REPEAT 750 1022 II.
 FT REPEAT 1200 1514 III.
 FT REPEAT 1523 1821 IV.
 FT TRANSMEM 124 147 S1 of repeat I.
 FT TRANSMEM 156 175 S2 of repeat I.
 FT TRANSMEM 189 207 S3 of repeat I.
 FT TRANSMEM 214 233 S4 of repeat I.
 FT TRANSMEM 250 273 S5 of repeat I.
 FT TRANSMEM 400 425 S6 of repeat I.
 FT TRANSMEM 763 787 S1 of repeat II.
 FT TRANSMEM 799 822 S2 of repeat II.
 FT TRANSMEM 831 850 S3 of repeat II.
 FT TRANSMEM 857 876 S4 of repeat II.
 FT TRANSMEM 893 913 S5 of repeat II.
 FT TRANSMEM 967 992 S6 of repeat II.
 FT TRANSMEM 1214 1237 S1 of repeat III.
 FT TRANSMEM 1251 1276 S2 of repeat III.
 FT TRANSMEM 1283 1304 S3 of repeat III.
 FT TRANSMEM 1309 1330 S4 of repeat III.
 FT TRANSMEM 1350 1377 S5 of repeat III.
 FT TRANSMEM 1457 1483 S6 of repeat III.
 FT TRANSMEM 1537 1560 S1 of repeat IV.
 FT TRANSMEM 1572 1595 S2 of repeat IV.
 FT TRANSMEM 1602 1625 S3 of repeat IV.
 FT TRANSMEM 1636 1657 S4 of repeat IV.
 FT TRANSMEM 1673 1695 S5 of repeat IV.
 FT TRANSMEM 1762 1786 S6 of repeat IV.
 FT CARBOHYD 211 211 N-linked (GlcNAc...)
 FT CARBOHYD 284 284 N-linked (GlcNAc...)
 FT CARBOHYD 295 295 N-linked (GlcNAc...)
 FT CARBOHYD 301 301 N-linked (GlcNAc...)
 FT CARBOHYD 306 306 N-linked (GlcNAc...)
 FT CARBOHYD 338 338 N-linked (GlcNAc...)
 FT CARBOHYD 601 601 N-linked (GlcNAc...)
 FT CARBOHYD 621 621 N-linked (GlcNAc...)
 FT CARBOHYD 681 681 N-linked (GlcNAc...)
 FT CARBOHYD 882 882 N-linked (GlcNAc...)
 FT CARBOHYD 1060 1060 N-linked (GlcNAc...)
 FT CARBOHYD 1064 1064 N-linked (GlcNAc...)
 FT CARBOHYD 1080 1080 N-linked (GlcNAc...)
 FT CARBOHYD 1146 1146 N-linked (GlcNAc...)
 FT CARBOHYD 1378 1378 N-linked (GlcNAc...)
 FT CARBOHYD 1392 1392 N-linked (GlcNAc...)
 FT CARBOHYD 1403 1403 N-linked (GlcNAc...)
 SQ SEQUENCE 2009 AA; 228769 MW; 680846f6368373B CRC64;

Query Match 100.0%; Score 28; DB 1; Length 2009;
 Best Local Similarity 100.0%; Pred. No. 6-7e-19;
 Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GIFFFVSYIIISFLVYVNNYIVILLENF 28
 |||||
 DB 1762 GIFFFVSYIIISFLVYVNNYIVILLENF 1769

RESULT 13
 O8CAJ8 PRELIMINARY; PRT; 253 AA.
 AC O8CAJ8; O8CAJ8;

DT 01-MAR-2003 (TrEMBLrel. 23, Created)
 DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
 DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
 DE Mus musculus adult male hypothalamus cDNA, RIKEN full-length enriched
 DE library, clone:A230052E19 product:sodium channel protein II homolog
 DE (fragment).
 GN Name=A230052E19R1k;
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Hypothalamus;
 RX MEDLINE=99279253; PubMed=10349636;
 RA Carninci P., Hayashizaki Y.;
 RT "High-efficiency full-length cDNA cloning";
 RL Meth. Enzymol. 303:19-44(1999).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Hypothalamus;
 RX MEDLINE=21085660; PubMed=11217851;
 RA RIKEN PANTOM Consortium;
 RT "Functional annotation of a full-length mouse cDNA collection.";
 RL Nature 409:685-690(2001).
 RN [3]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Hypothalamus;
 RA The PANTOM Consortium;
 RA The RIKEN Genome Exploration Research Group Phase I & II Team;
 RT "Analysis of the mouse transcriptome based on functional annotation of
 RT 60,770 full-length cDNAs";
 RL Nature 420:563-573(2002).
 RN [4]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Hypothalamus;
 RX MEDLINE=20499374; PubMed=11042159;
 RA Carninci P., Shibata Y., Hayatsu N., Sugahara Y., Shibata K., Itoh M.,
 RA Kono H., Okazaki Y., Muramatsu M., Hayashizaki Y.;
 RT "Normalization and subtraction of cap-trapper-selected cDNAs to
 RT prepare full-length cDNA libraries for rapid discovery of new genes.";
 RL Genome Res. 10:1617-1630(2000).
 RN [5]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Hypothalamus;
 RX MEDLINE=20530913; PubMed=11078681;
 RA Shibata K., Itoh M., Aizawa K., Nagaoka S., Sasaki N., Carninci P.,
 RA Kono H., Akiyama J., Nishi K., Katsunai T., Taishiro H., Itoh M.,
 RA Sumi N., Ishii Y., Nakamura S., Hazama M., Nishine T., Harada A.,
 RA Yamamoto R., Matsumoto H., Sakaguchi S., Ikegami T., Kashiwagi K.,
 RA Fujiwake S., Inoue K., Togawa Y., Izawa M., Ohara E., Matsuoka M.,
 RA Yoneda Y., Ishikawa T., Ozawa K., Tanaka T., Matsura S., Kawai J.,
 RA Okazaki Y., Muramatsu M., Inoue Y., Kira A., Hayashizaki Y.;
 RT "RIKEN integrated sequence analysis (RISA) system-384-format
 RT sequencing pipeline with 384 multicapillary sequencer.";
 RL Genome Res. 10:1157-1171(2000).
 RN [6]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Hypothalamus;
 RA Adachi J., Aizawa K., Akimura T., Arakawa T., Bono H., Carninci P.,
 RA Fukuda S., Furuno M., Hanagaki T., Hara A., Hashizume W.,
 RA Hayashida K., Hayatsu N., Hiramoto K., Hiraka T., Hirozane T.,
 RA Horii F., Imetani K., Ishii Y., Itoh M., Kagawa I., Kaubawa T.,
 RA Katoh H., Kawai U., Kojima Y., Kondo S., Kono H., Kouda M., Koya S.,
 RA Kurihara C., Matsuyama T., Miyazaki A., Murata M., Nakamura M.,
 RA Nishi K., Nomura K., Numazaki R., Ohno M., Ohsato N., Okazaki Y.,
 RA Saito R., Saitoh H., Sakai C., Sakai K., Sakazume N., Sano H.,
 RA Sasaki D., Shibata K., Shinagawa A., Shiraki T., Sogabe Y., Tagami M.,
 RA Tagawa A., Takahashi F., Takaku-Akahira S., Takeda Y., Tanaka T.,
 RA Tomaru A., Toya T., Yasunishi A., Muramatsu M., Hayashizaki Y.;
 RA Submitted (Jul-2001) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AK038645; BAC30078.1; -;
 DR MGD; MGI:2444703; A230052E19R1k.

DR GO: GO:0016020; C:membrane; IEA.
 DR GO: GO:0001518; C:voltage-gated sodium channel complex; IEA.
 DR GO: GO:0005261; F:cation channel activity; IEA.
 DR GO: GO:0005248; F:voltage-gated sodium channel activity; IEA.
 DR GO: GO:0006812; P:cation transport; IEA.
 DR GO: GO:0006814; P:sodium ion transport; IEA.
 DR InterPro: IPR001682; Ca/Na_pore.
 DR InterPro: IPR000048; IQ_region.
 DR Pfam: PF00612; IQ_1.
 DR PRINTS: PR00170; NACHANNEL.
 DR SMART: SM00015; IQ_1.
 DR PROSITE: PSS0096; IQ_1.
 KW Ionic channel.
 FT NON_TER 1
 SQ SEQUENCE 253 AA; 29041 MW; B6B1C4CC35A5E571E CRC64;

Query Match 96.4%; Score 27; DB 2; Length 253;
 Best Local Similarity 100.0%; Pred. No. 1.3e-18;
 Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 IFPVSYIIISFLVVMNYIAVILENF 28
 DB 1 IFPVSYIIISFLVVMNYIAVILENF 27

RESULT 14
 062205 PRELIMINARY; PRT; 309 AA.
 AC 062205;
 DT 01-NOV-1996 (TREMBLrel. 01, Created)
 DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
 DT 01-JUN-2003 (TREMBLrel. 24, Last annotation update)
 DE Sodium channel 25.
 GN Name=Scn9a;
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Brain;
 RA Jover E., Shah V.;
 RL Submitted (May-1995) to the EMBL/GenBank/DBJ databases.
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
 CC -1- SIMILARITY: Belongs to the sodium channel family.
 DR EMBL; L42338; AAA67106.1; -.
 DR MGD; MGI:107636; Scn9a.
 DR GO: GO:0016021; C:integral to membrane; IEA.
 DR GO: GO:0001518; C:voltage-gated sodium channel complex; IEA.
 DR GO: GO:0005261; F:cation channel activity; IEA.
 DR GO: GO:0005248; F:voltage-gated sodium channel activity; IEA.
 DR GO: GO:0006812; P:cation transport; IEA.
 DR GO: GO:0006814; P:sodium ion transport; IEA.
 DR InterPro: IPR001682; Ca/Na_pore.
 DR InterPro: IPR005821; Ion_trans.
 DR InterPro: IPR000048; IQ_region.
 DR InterPro: IPR005820; M+channel_nlg.
 DR InterPro: IPR001696; Na_channel.
 DR Pfam: PF00520; Ion_trans; 1.
 DR Pfam: PF00612; IQ_1.
 DR PRINTS: PR00170; NACHANNEL.
 DR SMART: SM00015; IQ_1.
 KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
 KW Transport; Voltage-gated channel.
 SQ SEQUENCE 309 AA; 34971 MW; 975165758B88D3EC CRC64;

Query Match 85.7%; Score 24; DB 2; Length 309;
 Best Local Similarity 100.0%; Pred. No. 1.4e-15;
 Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 5 FVSYIIISFLVVMNYIAVILENF 28
 DB 1 FVSYIIISFLVVMNYIAVILENF 28

DB 63 FVSYIIISFLVVMNYIAVILENF 86

RESULT 15
 015858 PRELIMINARY; PRT; 1977 AA.
 AC 015858;
 DT 01-NOV-1996 (TREMBLrel. 01, Created)
 DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
 DT 01-MAR-2004 (TREMBLrel. 26, Last annotation update)
 DE Sodium channel alpha subunit.
 GN Name=hNF-Na;
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Thyroid;
 RA MEDLINE=95237189; PubMed=7720699;
 RA Klugbauer N., Lacinova L., Flockezy V., Hofmann F.;
 RT "Structure and functional expression of a new member of the
 RT tetrodotoxin-sensitive voltage-activated sodium channel family from
 RT human neuroendocrine cells";
 RL EMBO J. 14:1084-1090(1995).
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
 CC -1- SIMILARITY: Belongs to the sodium channel family.
 DR EMBL; X82835; CAAS8042.1; -.
 DR PIR; S54771; S54771.
 DR HSSP; P04775; 1BYX.
 DR Genew; HGNC:10597; SCN9A.
 DR GO: GO:0005248; F:voltage-gated sodium channel activity; TAS.
 DR GO: GO:0006814; P:sodium ion transport; TAS.
 DR InterPro: IPR001682; Ca/Na_pore.
 DR InterPro: IPR002111; Cat_channel_tripL.
 DR InterPro: IPR005821; Ion_trans.
 DR InterPro: IPR000048; IQ_region.
 DR InterPro: IPR005820; M+channel_nlg.
 DR InterPro: IPR001696; Na_channel.
 DR InterPro: IPR010526; Na_trans_assoc.
 DR Pfam: PF00520; Ion_trans; 4.
 DR Pfam: PF00612; IQ_1.
 DR Pfam: PF06512; Na_trans_assoc; 1.
 DR PRINTS: PR00170; NACHANNEL.
 DR SMART: SM00015; IQ_1.
 KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
 KW Transport; Voltage-gated channel.
 SQ SEQUENCE 1977 AA; 225195 MW; 17D67C8C32BC15FB CRC64;

Query Match 85.7%; Score 24; DB 2; Length 1977;
 Best Local Similarity 100.0%; Pred. No. 5.7e-15;
 Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 5 FVSYIIISFLVVMNYIAVILENF 28
 DB 1729 FVSYIIISFLVVMNYIAVILENF 1752

RESULT 16
 028644 PRELIMINARY; PRT; 1984 AA.
 AC 028644;
 DT 01-NOV-1996 (TREMBLrel. 01, Created)
 DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
 DT 01-MAR-2004 (TREMBLrel. 26, Last annotation update)
 DE Sodium channel alpha-subunit.
 GN Oryctolagus cuniculus (Rabbit).
 OS Oryctolagus cuniculus (Rabbit).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Lagomorpha; Leporidae; Oryctolagus.
 OX NCBI_TaxID=9986;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=New Zealand White; TISSUE=Sciatic nerve;

RX MEDLINE=96074641; PubMed=7479931;
 RA Belcher S.M., Zerillo C.A., Levenson R., Ritchie J.M., Howe J.R.;
 RT "Cloning of a sodium channel alpha subunit from rabbit Schwann
 cells."
 RL Proc. Natl. Acad. Sci. U.S.A. 92:11034-11038(1995).
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
 DR EMBL; U55238; AAB89159.1; -.
 DR HSSP; P04775; 1BYV.
 DR GO; GO:0016021; C:integral to membrane; IEA.
 DR GO; GO:0001515; C:voltage-gated sodium channel complex; IEA.
 DR GO; GO:0005261; F:cation channel activity; IEA.
 DR GO; GO:0005248; F:voltage-gated sodium channel activity; IEA.
 DR GO; GO:0006812; P:cation transport; IEA.
 DR GO; GO:0006814; P:sodium ion transport; IEA.
 DR InterPro; IPR001682; Ca/Na_pore.
 DR InterPro; IPR002111; Cat_channel_TrypL.
 DR InterPro; IPR005821; Ion_trans.
 DR InterPro; IPR000048; IQ_region.
 DR InterPro; IPR005820; M+channel_nlg.
 DR InterPro; IPR001696; Na_channel_nlg.
 DR InterPro; IPR010526; Na_trans_assoc.
 DR Pfam; PF00520; Ion_trans; 4.
 DR Pfam; PF00612; IQ_1.
 DR Pfam; PF06512; Na_trans_assoc; 1.
 DR PRINTS; PR00170; NACHANNEL.
 DR SMART; SM00015; IQ_1.
 DR Ion transport; Ionic channel; Sodium channel; Transmembrane;
 KM Transport; Voltage-gated channel.
 SQ SEQUENCE 1984 AA; 225748 MW; 98F76860C9866AA0 CRC64;

Query Match 85.7%; Score 24; DB 2; Length 1984;
 Best Local Similarity 100.0%; Pred. No. 5.8e-15;
 Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 FVSYIIISFLVVMYIAVILENF 28
 Db 1737 FVSYIIISFLVVMYIAVILENF 1760

RESULT 17
 ID 008562 PRELIMINARY; PRT; 1984 AA.
 AC 008562;
 DT 01-JUL-1997 (TrEMBLrel. 04, Created)
 DT 01-JUL-1997 (TrEMBLrel. 04, Last sequence update)
 DT 05-JUL-2004 (TrEMBLrel. 27, Last annotation update)
 DE PNI (Voltage-gated sodium channel) (Fragment).
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 OC NCBI_TaxID=10116;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=97188502; PubMed=9037087;
 RA Toledo-Aral J.J., Mose B.L., He Z.J., Kozowski A.G., Whisenand T.,
 Levinson S.R., Wolf J.J., Silos-Santiago I., Halesova S., Mandel G.;
 RT "Identification of PNI, a predominant voltage-dependent sodium channel
 expressed principally in peripheral neurons."
 RL Proc. Natl. Acad. Sci. U.S.A. 94:1527-1532(1997).
 RN [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=97007982; PubMed=8654872;
 RA Kozak C.A., Sangameswaran L.;
 RT "Genetic mapping of the peripheral sodium channel genes, Scn5a and
 Scn10a, in the mouse."
 RL Mamm. Genome 7:787-788(1996).
 RN [3]
 RP SEQUENCE FROM N.A.
 RA Sangameswaran L., Fish L.M., Koch B.D., Rabert D.K., Delgado S.G.,
 RA Ilinck M., Jakeman L.B., Novakovic S., Wong K., Sze P., Tzoumaka E.,
 RA Stewart G.R., Herman R.C., Chan H., Eglen R.M., Hunter J.C.;
 RT "A novel tetrodotoxin-sensitive, voltage-gated sodium channel

RT expressed in rat and human dorsal root ganglia."
 RL J. Biol. Chem. 0:0-0(1997).
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
 DR EMBL; U79568; AAB50403.1; -.
 DR EMBL; AF000368; AAB80701.1; -.
 DR HSSP; P04775; 1BYV.
 DR GO; GO:0016021; C:integral to membrane; IEA.
 DR GO; GO:0001515; C:voltage-gated sodium channel complex; IEA.
 DR GO; GO:0005261; F:cation channel activity; IEA.
 DR GO; GO:0005248; F:voltage-gated sodium channel activity; IEA.
 DR GO; GO:0006812; P:cation transport; IEA.
 DR GO; GO:0006814; P:sodium ion transport; IEA.
 DR InterPro; IPR001682; Ca/Na_pore.
 DR InterPro; IPR002111; Cat_channel_TrypL.
 DR InterPro; IPR005821; Ion_trans.
 DR InterPro; IPR000048; IQ_region.
 DR InterPro; IPR005820; M+channel_nlg.
 DR InterPro; IPR001696; Na_channel.
 DR InterPro; IPR010526; Na_trans_assoc.
 DR Pfam; PF00520; Ion_trans; 4.
 DR Pfam; PF00612; IQ_1.
 DR Pfam; PF06512; Na_trans_assoc; 1.
 DR PRINTS; PR00170; NACHANNEL.
 DR SMART; SM00015; IQ_1.
 DR Ion transport; Ionic channel; Sodium channel; Transmembrane;
 KM Transport; Voltage-gated channel.
 FT NON_TER
 SQ SEQUENCE 1984 AA; 226037 MW; 386C389B5097091 CRC64;

Query Match 85.7%; Score 24; DB 2; Length 1984;
 Best Local Similarity 100.0%; Pred. No. 5.8e-15;
 Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 FVSYIIISFLVVMYIAVILENF 28
 Db 1738 FVSYIIISFLVVMYIAVILENF 1761

RESULT 18
 ID 054811 PRELIMINARY; PRT; 279 AA.
 AC 054811;
 DT 01-JUN-1998 (TrEMBLrel. 06, Created)
 DT 01-JUN-1998 (TrEMBLrel. 06, Last sequence update)
 DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
 DE GPBI (Fragment).
 OS Cavia porcellus (Guinea pig).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Hystriocognathi; Caviidae; Cavia.
 OC NCBI_TaxID=10141;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=97338144; PubMed=9192691;
 RA de Miera E.V.S., Rudy B., Sugimori M., Ilinas R.;
 RT "Molecular characterization of the sodium channel subunits expressed
 in mammalian cerebellar Purkinje cells."
 RL Proc. Natl. Acad. Sci. U.S.A. 94:7059-7064(1997).
 RN [2]
 RP SEQUENCE FROM N.A.
 RX TISSUE=Cerebellum;
 RA Vega-Saenz de Miera E., Rudy B., Sugimori M., Ilinas R.;
 RL Submitted (May-1997) to the EMBL/GenBank/DBD databases.
 DR EMBL; AF003372; AAC02899.1; -.
 DR GO; GO:0016021; C:integral to membrane; IEA.
 DR GO; GO:0008076; C:voltage-gated potassium channel complex; IEA.
 DR GO; GO:0005261; F:cation channel activity; IEA.
 DR GO; GO:0005249; F:voltage-gated potassium channel activity; IEA.
 DR GO; GO:0006812; P:cation transport; IEA.
 DR GO; GO:0006814; P:potassium ion transport; IEA.
 DR InterPro; IPR001682; Ca/Na_pore.
 DR InterPro; IPR005821; Ion_trans.

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DR InterPro; IPR003091; K_channel.
DR InterPro; IPR005820; M+channel_nlg.
DR Pfam; PF00520; Ion_trans; 1.
DR PRINTS; PR00169; KCHANNEL.
KW Ion transport; Ionic channel; Transmembrane; Transport.
FT NON_TER 279 279
SQ SEQUENCE 279 AA; 31625 MW; 1504A333CA63DD24 CRC64;

Query Match 60.7%; Score 17; DB 2; Length 279;
Best Local Similarity 100.0%; Pred. No. 1e-08;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GIFFPVSYIIISFLVV 17
DB 263 GIFFVSYIIISFLVV 279

RESULT 19
P0670 PRELIMINARY; PRT; 1993 AA.
ID P0670;
AC P0670;
DT 01-MAY-1997 (TrEMBLrel. 03, Created)
DT 01-MAY-1997 (TrEMBLrel. 03, Last sequence update)
DE 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE Sodium channel alpha-subunit SCAP1.
OS Aplysia californica (California sea hare).
OC Eukaryota; Metazoa; Mollusca; Gastropoda; Orthogastropoda;
OC Apogastropoda; Heterobranchia; Euthyneura; Opisthobranchia; Anaspidea;
OC Aplysioidae; Aplysiidae; Aplysia.
OX NCBI_Taxid=6500;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Nervous system;
RX MEDLINE=97238630; PubMed=9115644;
RA Dyer J.R., Johnston W.L., Castellucci V.F., Dunn R.J.;
RT "Cloning and tissue distribution of the Aplysia Na+ channel alpha-
RT subunit cDNA."
RL DNA Cell Biol. 16:347-356(1997).
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
CC -1- SIMILARITY: Belongs to the sodium channel family.
DR EMBL; U66915; AAC47457.1; -.
DR PIR; T30902; T30902.
DR HSSP; P04775; 1BY.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0001518; C:voltage-gated sodium channel complex; IEA.
DR GO; GO:0005261; F:cation channel activity; IEA.
DR GO; GO:0005248; F:voltage-gated sodium channel activity; IEA.
DR GO; GO:0006812; P:cation transport; IEA.
DR GO; GO:0006814; P:sodium ion transport; IEA.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR002111; Cat_channel_TrpL.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR005820; M+channel_nlg.
DR InterPro; IPR001696; Na_channel.
DR InterPro; IPR010526; Na_trans_assoc.
DR Pfam; PF00520; Ion_trans; 4.
DR Pfam; PF06512; Na_trans_assoc; 1.
DR PRINTS; PR00170; NACHANNEL.
KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
KW Transport; Voltage-gated channel.
SQ SEQUENCE 1993 AA; 225896 MW; 33E174B9BF07E1A7 CRC64;

Query Match 60.7%; Score 17; DB 2; Length 1993;
Best Local Similarity 100.0%; Pred. No. 4.5e-08;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 12 SFLVVNMVYAVILENF 28
DB 1719 SFLVVNMVYAVILENF 1735

RESULT 20
Q9IBF1

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ID Q9IBF1 PRELIMINARY; PRT; 1880 AA.
AC Q9IBF1;
DT 01-OCT-2000 (TrEMBLrel. 15, Created)
DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE Voltage-gated sodium channel.
DE Takifugu pardalis (Puffer puffer).
OS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
OC Acanthomorpha; Acanthopterygii; Percomorphi; Tetraodontiformes;
OC Tetraodontidae; Tetraodontidae; Takifugu.
OX NCBI_Taxid=98921;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=skeletal muscle;
RX MEDLINE=20090650; PubMed=10623632;
RA Yotsu-Yamashita M., Nishimori K., Nitanai Y., Isemura M., Sugimoto A.,
RA Yasumoto T.;
RT "Binding properties of 3H-PbTx-3 and 3H-saxitoxin to brain membranes
RT and to skeletal muscle membranes of puffer fish Takifugu pardalis, and the
RT primary structure of a voltage-gated Na+ channel alpha-subunit (fknal)
RL from skeletal muscle of F. pardalis."
RL Biochem. Biophys. Res. Commun. 267:403-412(2000).
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
CC -1- SIMILARITY: Belongs to the sodium channel family.
DR EMBL; AB030482; BAA90398.1; -.
DR HSSP; P04775; 1BY.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0001518; C:voltage-gated sodium channel complex; IEA.
DR GO; GO:0005261; F:cation channel activity; IEA.
DR GO; GO:0005248; F:voltage-gated sodium channel activity; IEA.
DR GO; GO:0006812; P:cation transport; IEA.
DR GO; GO:0006814; P:sodium ion transport; IEA.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR002111; Cat_channel_TrpL.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IQ_region.
DR InterPro; IPR005820; M+channel_nlg.
DR InterPro; IPR001696; Na_channel.
DR InterPro; IPR010526; Na_trans_assoc.
DR Pfam; PF00520; Ion_trans; 4.
DR Pfam; PF00612; IQ; 1.
DR Pfam; PF06512; Na_trans_assoc; 1.
DR PRINTS; PR00170; NACHANNEL.
DR PROSITE; PS50096; IQ; 1.
KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
KW Transport; Voltage-gated channel.
SQ SEQUENCE 1880 AA; 212084 MW; 406483C6C3D43E02 CRC64;

Query Match 57.1%; Score 16; DB 2; Length 1880;
Best Local Similarity 100.0%; Pred. No. 4.2e-07;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 7 SYIIISFLVVNMVYIA 22
DB 1608 SYIIISFLVVNMVYIA 1623

RESULT 21
OS4812 PRELIMINARY; PRT; 278 AA.
ID OS4812;
AC OS4812;
DT 01-JUN-1998 (TrEMBLrel. 06, Created)
DT 01-JUN-1998 (TrEMBLrel. 06, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE Ceriiti (Fragment).
OS Cavia porcellus (Guinea pig).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Hystricognathi; Caviidae; Cavia.
OX NCBI_Taxid=10141;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Cerebellum;

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RX MEDLINE=97338144; PubMed=9192691;
RA de Miera E.V.S., Rudy B., Sugimori M., Llinas R.;
RT "Molecular characterization of the sodium channel subunits expressed
RT in mammalian cerebellar Purkinje cells";
RT Proc. Natl. Acad. Sci. U.S.A. 94:7059-7064(1997).
RN (12)
RN SEQUENCE FROM N.A.
RP TISSUE=Cerebellum;
RA Vega-Saenz de Miera E., Rudy B., Sugimori M., Llinas R.;
RL Submitted (MAY-1997) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF003373; AAC02900.1; -.
DR GO; GO:0016021; C:Integral to membrane; IEA.
DR GO; GO:0008076; C:voltage-gated potassium channel complex; IEA.
DR GO; GO:0005261; P:cation channel activity; IEA.
DR GO; GO:0005249; P:voltage-gated potassium channel activity; IEA.
DR GO; GO:0006812; P:cation transport; IEA.
DR GO; GO:0006813; P:potassium ion transport; IEA.
DR InterPro; IPR001682; Ca/Na_pore..
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR003091; K_channel.
DR InterPro; IPR005820; M-channel_nlg.
DR Pfam; PF00520; Ion_trans; 1.
DR PRINTS; PR00169; KCHANNEL.
DR ION_TRANSPORT; Ionic channel; Transmembrane; Transport.
DR NON_TER 278
FT SEQUENCE 278 AA; 31866 MW; 65741B14C6A649A CRC64;
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Best Local Similarity 100.0%; Pred. No. 9.2e-06;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GIFFVSYIIISFL 14
DB 262 GIFFVSYIIISFL 275
RESULT 22
063360 PRELIMINARY; PRT; 324 AA.
AC 063360;
DT 01-NOV-1996 (TREMBLrel. 01, Created)
DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
DT 01-JUN-2003 (TREMBLrel. 24, Last annotation update)
DE Na+ channel (Fragment).
CN Name=Na+ channel;
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OC NCBI_TaxId=10116;
OX (1)
RN SEQUENCE FROM N.A.
RP TISSUE=Myocardium;
RA MEDLINE=89292178; PubMed=2544627;
RA Sills M.N., Xu Y.C., Baracchini E., Goodman R.H., Cooperman S.S.,
RA Mandel G., Chien K.R.;
RT "Expression of diverse Na+ channel messenger RNAs in rat myocardium:
RT Evidence for a cardiac-specific Na+ channel.";
RT J. Clin. Invest. 84:331-336(1989).
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
CC -1- SIMILARITY: Belongs to the sodium channel family.
DR EMBL; M27223; AAA1666.1; -.
DR PIR; A45752; A45752.
DR GO; GO:0016021; C:Integral to membrane; IEA.
DR GO; GO:0001518; C:voltage-gated sodium channel complex; IEA.
DR GO; GO:0005261; P:cation channel activity; IEA.
DR GO; GO:0005248; P:voltage-gated sodium channel activity; IEA.
DR GO; GO:0006812; P:cation transport; IEA.
DR GO; GO:0006814; P:sodium ion transport; IEA.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR002111; Cat_channel_TpPL.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IO_region.
DR Pfam; PF00612; IQ; 1.
DR PRINTS; PR00170; NACHANNEL.
DR SMART; SM00015; IQ; 1.
DR PROSITE; PS50096; IQ; 1.
DR ION_TRANSPORT; Ionic channel; Sodium channel; Transmembrane;
DR Transport; Voltage-gated channel.
FT NON_TER 364
FT SEQUENCE 364 AA; 41263 MW; BA5760C962BE6786 CRC64;
SQ
Query Match 50.0%; Score 14; DB 2; Length 364;
Best Local Similarity 100.0%; Pred. No. 1.1e-05;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GIFFVSYIIISFL 14
DB 131 GIFFVSYIIISFL 144
RESULT 24

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DR InterPro; IPR001696; Na_channel.
DR Pfam; PF00520; Ion_trans; 1.
DR Pfam; PF00612; IQ; 1.
DR PRINTS; PR00170; NACHANNEL.
DR SMART; SM00015; IQ; 1.
DR PROSITE; PS50096; IQ; 1.
DR ION_TRANSPORT; Ionic channel; Sodium channel; Transmembrane;
DR Transport; Voltage-gated channel.
FT NON_TER 1
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SQ
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Best Local Similarity 100.0%; Pred. No. 1e-05;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GIFFVSYIIISFL 14
DB 86 GIFFVSYIIISFL 99
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ID 09P206;
AC 09P206;
DT 01-OCT-2000 (TREMBLrel. 15, Created)
DT 01-OCT-2000 (TREMBLrel. 15, Last sequence update)
DT 01-JUN-2003 (TREMBLrel. 24, Last annotation update)
DE Voltage-gated sodium channel alpha subunit (Fragment).
GN Name=SCN8A;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OC NCBI_TaxId=9606;
OX (1)
RN SEQUENCE FROM N.A.
RP Kitanu T., Kobayakawa H., Saitou N.;
RA Submitted (JAN-2000) to the EMBL/GenBank/DBJ databases.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
CC -1- SIMILARITY: Belongs to the sodium channel family.
DR EMBL; AB037525; BA90445.1; -.
DR GO; GO:0016021; C:Integral to membrane; IEA.
DR GO; GO:0001518; C:voltage-gated sodium channel complex; IEA.
DR GO; GO:0005261; P:cation channel activity; IEA.
DR GO; GO:0005248; P:voltage-gated sodium channel activity; IEA.
DR GO; GO:0006812; P:cation transport; IEA.
DR GO; GO:0006814; P:sodium ion transport; IEA.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR002111; Cat_channel_TpPL.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IO_region.
DR InterPro; IPR005820; M-channel_nlg.
DR InterPro; IPR001696; Na_channel.
DR Pfam; PF00520; Ion_trans; 1.
DR Pfam; PF00612; IQ; 1.
DR PRINTS; PR00170; NACHANNEL.
DR SMART; SM00015; IQ; 1.
DR PROSITE; PS50096; IQ; 1.
DR ION_TRANSPORT; Ionic channel; Sodium channel; Transmembrane;
DR Transport; Voltage-gated channel.
FT NON_TER 364
FT SEQUENCE 364 AA; 41263 MW; BA5760C962BE6786 CRC64;
SQ
Query Match 50.0%; Score 14; DB 2; Length 364;
Best Local Similarity 100.0%; Pred. No. 1.1e-05;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GIFFVSYIIISFL 14
DB 131 GIFFVSYIIISFL 144
RESULT 24

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Q9N2E3
ID Q9N2E3 PRELIMINARY; PRT; 364 AA.
AC Q9N2E3;
DT 01-OCT-2000 (TRENBLrel. 15, Created)
DT 01-OCT-2000 (TRENBLrel. 15, Last sequence update)
DT 01-JUN-2003 (TRENBLrel. 24, Last annotation update)
DE Voltage-gated sodium channel alpha subunit (Fragment).
GN Name=SCN8A;
OS Pongo pygmaeus (Orangutan).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Pongo.
OX NCBI_TaxID=9600;
RN [1]
RP SEQUENCE FROM N.A.
RA Kitano T., Kobayakawa H., Saitou N.;
RL Submitted (JAN-2000) to the EMBL/GenBank/DBJ databases.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
CC -1- SIMILARITY: Belongs to the sodium channel family.
DR EMBL; AB037528; BAA90448.1; -.
DR GO; GO:0016021; C:Integral to membrane; IEA.
DR GO; GO:0001518; C:voltage-gated sodium channel complex; IEA.
DR GO; GO:0005248; F:cation channel activity; IEA.
DR GO; GO:0006812; P:voltage-gated sodium channel activity; IEA.
DR GO; GO:0006814; P:sodium ion transport; IEA.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR002111; Cat_channel_TrpL.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IQ_region.
DR InterPro; IPR005820; M_channel_nlg.
DR Pfam; PF00520; Ion_trans; 1.
DR PRINTS; PR00170; NACHANNEL.
DR SMART; SM00015; IQ; 1.
DR PROSITE; PS50096; IQ; 1.
KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
KW Transport; Voltage-gated channel.
FT NON_TER 1
FT NON_TER 364
SQ SEQUENCE 364 AA; 41235 MW; 975760C962BE6791 CRC64;

Query Match 50.0%; Score 14; DB 2; Length 364;
Best Local Similarity 100.0%; Pred. No. 1.1e-05;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GIFFVSYIIISFL 14
Db 131 GIFFVSYIIISFL 144

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RESULT 25
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AC Q9N2E4;
DT 01-OCT-2000 (TRENBLrel. 15, Created)
DT 01-OCT-2000 (TRENBLrel. 15, Last sequence update)
DT 01-JUN-2003 (TRENBLrel. 24, Last annotation update)
DE Voltage-gated sodium channel alpha subunit (Fragment).
GN Name=SCN8A;
OS Gorilla gorilla (gorilla).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Gorilla.
OX NCBI_TaxID=9593;
RN [1]
RP SEQUENCE FROM N.A.
RA Kitano T., Kobayakawa H., Saitou N.;
RL Submitted (JAN-2000) to the EMBL/GenBank/DBJ databases.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (By similarity).
CC -1- SIMILARITY: Belongs to the sodium channel family.
DR EMBL; AB037527; BAA90447.1; -.
DR GO; GO:0016021; C:Integral to membrane; IEA.
DR GO; GO:0006812; P:voltage-gated sodium channel complex; IEA.

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DR GO; GO:0005261; F:cation channel activity; IEA.
DR GO; GO:0005248; F:voltage-gated sodium channel activity; IEA.
DR GO; GO:0006812; P:cation transport; IEA.
DR GO; GO:0006814; P:sodium ion transport; IEA.
DR InterPro; IPR001682; Ca/Na_pore.
DR InterPro; IPR002111; Cat_channel_TrpL.
DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR000048; IQ_region.
DR InterPro; IPR005820; M_channel_nlg.
DR InterPro; IPR001696; Na_channel.
DR Pfam; PF00520; Ion_trans; 1.
DR PRINTS; PR00170; NACHANNEL.
DR PROSITE; PS50096; IQ; 1.
KW Ion transport; Ionic channel; Sodium channel; Transmembrane;
KW Transport; Voltage-gated channel.
FT NON_TER 1
FT NON_TER 364
SQ SEQUENCE 364 AA; 41263 MW; BA5760C962BE6786 CRC64;

Query Match 50.0%; Score 14; DB 2; Length 364;
Best Local Similarity 100.0%; Pred. No. 1.1e-05;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GIFFVSYIIISFL 14
Db 131 GIFFVSYIIISFL 144

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Search completed: January 27, 2005, 17:51:35
Job time : 94.5 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 27, 2005, 17:36:50 ; Search time 22.5 Seconds
(without alignments)
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Title: US-10-608-584-29

Perfect score: 28
Sequence: 1 GIFFFVXYIIISFLVVMYIAVLLENF 28

Scoring table: OLIGO
Gapop 60.0 , Gapext 60.0

Searched: 478139 seqs, 66318000 residues

Word size : 0

Total number of hits satisfying chosen parameters: 478139

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 100 summaries

Database :

Issued Patents AA: *
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Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

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78	6	21.4	120 4 US-09-252-991A-26987	Sequence 26987, A
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87	6	21.4	204 4 US-09-270-767-56679	Sequence 56679, A
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ALIGNMENTS

RESULT 1

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US-08-605-284B-13
; Sequence 13, Application US/08605284B
; Patent No. 6060271
; GENERAL INFORMATION:
; APPLICANT: WALEWSKI, JOSE L.
; TITLE OF INVENTION: VOLTAGE GATED SODIUM CHANNELS FROM
; TITLE OF INVENTION: HUMAN PERIPHERAL NERVE
; NUMBER OF SEQUENCES: 23
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: NIXON, HARGRAVE, DEVANS & DOYLE LLP
; STREET: CLINTON SQUARE, P.O. BOX 1051
; CITY: ROCHESTER
; STATE: NEW YORK
; COUNTRY: USA
; ZIP: 14603
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/605,284B
; FILING DATE: 09-FEB-1996
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: BRAMAN, SUSAN J.
; REGISTRATION NUMBER: 34,103
; REFERENCE/DOCKET NUMBER: 19603/800 (CRF D-1705)
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 716-263-1636
; TELEFAX: 716-263-1636
; INFORMATION FOR SEQ ID NO: 13:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 310 amino acids
; TYPE: amino acid
; STRANDEDNESS: not relevant
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-605-284B-13

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Query Match

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Best Local Similarity 100.0%; Score 28; DB 3; Length 310;
Pred. No. 5.1e-21;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 1 GIFFFVSYIIISFLVVMNYIAVILENF 28
DB 283 GIFFFVSYIIISFLVVMNYIAVILENF 310

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RESULT 2

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US-08-605-284B-14
; Sequence 14, Application US/08605284B
; Patent No. 6060271
; GENERAL INFORMATION:
; APPLICANT: WALEWSKI, JOSE L.
; TITLE OF INVENTION: VOLTAGE GATED SODIUM CHANNELS FROM
; TITLE OF INVENTION: HUMAN PERIPHERAL NERVE
; NUMBER OF SEQUENCES: 23
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: NIXON, HARGRAVE, DEVANS & DOYLE LLP
; STREET: CLINTON SQUARE, P.O. BOX 1051
; CITY: ROCHESTER
; STATE: NEW YORK
; COUNTRY: USA
; ZIP: 14603

```

COMPUTER READABLE FORM:

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; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/605,284B
; FILING DATE: 09-FEB-1996
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: BRAMAN, SUSAN J.
; REGISTRATION NUMBER: 34,103
; REFERENCE/DOCKET NUMBER: 19603/800 (CRF D-1705)
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 716-263-1636
; TELEFAX: 716-263-1636
; INFORMATION FOR SEQ ID NO: 14:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 310 amino acids
; TYPE: amino acid
; STRANDEDNESS: not relevant
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-605-284B-14

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Query Match

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Best Local Similarity 100.0%; Score 28; DB 3; Length 310;
Pred. No. 5.1e-21;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 1 GIFFFVSYIIISFLVVMNYIAVILENF 28
DB 283 GIFFFVSYIIISFLVVMNYIAVILENF 310

```

RESULT 3

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US-08-605-284B-15
; Sequence 15, Application US/08605284B
; Patent No. 6060271
; GENERAL INFORMATION:
; APPLICANT: WALEWSKI, JOSE L.
; TITLE OF INVENTION: VOLTAGE GATED SODIUM CHANNELS FROM
; TITLE OF INVENTION: HUMAN PERIPHERAL NERVE
; NUMBER OF SEQUENCES: 23
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: NIXON, HARGRAVE, DEVANS & DOYLE LLP
; STREET: CLINTON SQUARE, P.O. BOX 1051
; CITY: ROCHESTER
; STATE: NEW YORK
; COUNTRY: USA
; ZIP: 14603
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/605,284B
; FILING DATE: 09-FEB-1996
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: BRAMAN, SUSAN J.
; REGISTRATION NUMBER: 34,103
; REFERENCE/DOCKET NUMBER: 19603/800 (CRF D-1705)
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 716-263-1636
; TELEFAX: 716-263-1636
; INFORMATION FOR SEQ ID NO: 15:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 310 amino acids
; TYPE: amino acid
; STRANDEDNESS: not relevant
; TOPOLOGY: linear

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MOLECULE TYPE: protein
US-08-605-284B-15

Query Match 100.0%; Score 28; DB 3; Length 310;
Best Local Similarity 100.0%; Pred. No. 5.1e-21;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GIFFVSYIIISFLVVMNYIAVILENF 28
|||
Db 283 GIFFVSYIIISFLVVMNYIAVILENF 310

RESULT 4

US-08-836-325-7
; Sequence 7, Application US/08836325
; Patent No. 6110672

GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Halegoua, Simon
APPLICANT: Borden, Laurence A.
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
TITLE OF INVENTION: Thereof
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/836,325
FILING DATE: 2-MAY-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917,0240002
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 2005 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-08-836-325-7

Query Match 100.0%; Score 28; DB 3; Length 2005;
Best Local Similarity 100.0%; Pred. No. 2.9e-20;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GIFFVSYIIISFLVVMNYIAVILENF 28
|||
Db 1 GIFFVSYIIISFLVVMNYIAVILENF 28

Db 1752 GIFFVSYIIISFLVVMNYIAVILENF 1779

RESULT 5
US-09-457-571-7
; Sequence 7, Application US/09457571
; Patent No. 6703486

GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Halegoua, Simon
APPLICANT: Borden, Laurence A.
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
TITLE OF INVENTION: Thereof
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/457,571
FILING DATE: 09-DEC-1999
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/836,325
FILING DATE: 02-MAY-1997
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917,0240003
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 2005 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-457-571-7

Query Match 100.0%; Score 28; DB 4; Length 2005;
Best Local Similarity 100.0%; Pred. No. 2.9e-20;
Matches 28; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GIFFVSYIIISFLVVMNYIAVILENF 28
|||
Db 1752 GIFFVSYIIISFLVVMNYIAVILENF 1779

RESULT 6
US-08-605-284B-4
; Sequence 4, Application US/08605284B
; Patent No. 6060271

```

GENERAL INFORMATION:
APPLICANT: WALEMSKI, JOSE L.
TITLE OF INVENTION: VOLTAGE GATED SODIUM CHANNELS FROM
NUMBER OF SEQUENCES: 23
CORRESPONDENCE ADDRESS:
ADDRESS: NIXON, HARGRAVE, DEVANS & DOYLE LLP
CITY: ROCHESTER
STATE: NEW YORK
COUNTRY: USA
ZIP: 14603
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/605,284B
FILING DATE: 09-FEB-1996
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: BRAMAN, SUSAN J.
REGISTRATION NUMBER: 34,103
REFERENCE/DOCKET NUMBER: 19603/800 (CRF D-1705)
TELEPHONE: 716-263-1636
TELEFAX: 716-263-1600
INFORMATION FOR SEQ ID NO:
SEQUENCE CHARACTERISTICS:
LENGTH: 309 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-605-284B-4

Query Match
Best Local Similarity 96.4%; Score 27; DB 3; Length 309;
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 GIFFFVSIIISFLVVMNYIAVILEN 27
283 GIFFFVSIIISFLVVMNYIAVILEN 309

```

```

ATTORNEY/AGENT INFORMATION:
NAME: BRAMAN, SUSAN J.
REGISTRATION NUMBER: 34,103
REFERENCE/DOCKET NUMBER: 19603/800 (CRF D-1705)
TELEPHONE: 716-263-1636
TELEFAX: 716-263-1600
INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 309 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-605-284B-5

Query Match
Best Local Similarity 96.4%; Score 27; DB 3; Length 309;
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 GIFFFVSIIISFLVVMNYIAVILEN 27
283 GIFFFVSIIISFLVVMNYIAVILEN 309

RESULT 6
US-08-836-325-8
Sequence 8, Application US/08836325
Patent No. 6110672
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Halegoua, Simon
TITLE OF INVENTION: Borden, Laurence A.
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESS: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/836,325
FILING DATE: 2-MAY-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917, 0240002
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 8:
SEQUENCE CHARACTERISTICS:

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LENGTH: 813 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-08-836-325-8

Query Match 89.3%; Score 25; DB 3; Length 813;
Best Local Similarity 100.0%; Pred. No. 1.4e-17;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 FVSYIIISFLVVMYIAVILENF 28
|||||
DB 617 FVSYIIISFLVVMYIAVILENF 641

RESULT 9
US-09-457-571-8
Sequence 8, Application US/09457571
Patent No. 6703486

GENERAL INFORMATION:
APPLICANT: Mandel, Gail
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
TITLE OF INVENTION: Thereof
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C.
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/457,571
FILING DATE: 09-DEC-1999
CLASSIFICATION: 514

PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/836,325
FILING DATE: 02-MAY-1997
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917.0240003
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540

INFORMATION FOR SEQ ID NO: 8:
SEQUENCE CHARACTERISTICS:
LENGTH: 813 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-457-571-8

Query Match 89.3%; Score 25; DB 4; Length 813;
Best Local Similarity 100.0%; Pred. No. 1.4e-17;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 FVSYIIISFLVVMYIAVILENF 28
|||||
DB 617 FVSYIIISFLVVMYIAVILENF 641

RESULT 10
US-08-836-325-2
Sequence 2, Application US/08836325
Patent No. 6110672

GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Halegoua, Simon
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
TITLE OF INVENTION: Thereof
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C.
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/836,325
FILING DATE: 2-MAY-1997
CLASSIFICATION: 514

PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917.0240002
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540

INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 1011 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-836-325-2

Query Match 85.7%; Score 24; DB 3; Length 1011;
Best Local Similarity 100.0%; Pred. No. 1.8e-16;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 FVSYIIISFLVVMYIAVILENF 28
|||||
DB 785 FVSYIIISFLVVMYIAVILENF 808

RESULT 11

US-09-457-571-2
; Sequence 2, Application US/09457571
; Patent No. 6703486
; GENERAL INFORMATION:
; APPLICANT: Mandel, Gail
; APPLICANT: Halegoua, Simon
; TITLE OF INVENTION: Peripheral Nervous System Specific
; TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
; TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
; TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
; TITLE OF INVENTION: thereof
; NUMBER OF SEQUENCES: 19
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C
; STREET: 1100 New York Ave., N. W., Suite 600
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20005-3934
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/457,571
; FILING DATE: 09-DEC-1999
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/836,325
; FILING DATE: 02-MAY-1997
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/14251
; FILING DATE: 02-NOV-1995
; APPLICATION NUMBER: 08/482,401
; FILING DATE: 07-JUN-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/334,029
; FILING DATE: 02-NOV-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Ludwig, Steven R.
; REGISTRATION NUMBER: 36,203
; REFERENCE/DOCKET NUMBER: 0917.0240003
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-371-2600
; TELEFAX: 202-371-2540
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1011 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-09-457-571-2

Query Match 85.7%; Score 24; DB 4; Length 1011;
Best Local Similarity 100.0%; Pred. No. 18e-16;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 5 FVSYIIISFLVVMNYIAVILENF 28
785 FVSYIIISFLVVMNYIAVILENF 808

RESULT 12
US-08-836-325-15
; Sequence 15, Application US/08836325
; Patent No. 6110672
; GENERAL INFORMATION:
; APPLICANT: Mandel, Gail
; APPLICANT: Halegoua, Simon
; APPLICANT: Borden, Laurence A.
; TITLE OF INVENTION: Peripheral Nervous System Specific

; TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
; TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
; TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
; TITLE OF INVENTION: thereof
; NUMBER OF SEQUENCES: 19
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C
; STREET: 1100 New York Ave., N. W., Suite 600
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20005-3934
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/836,325
; FILING DATE: 2-MAY-1997
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/14251
; FILING DATE: 02-NOV-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/482,401
; FILING DATE: 07-JUN-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/334,029
; FILING DATE: 02-NOV-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Ludwig, Steven R.
; REGISTRATION NUMBER: 36,203
; REFERENCE/DOCKET NUMBER: 0917.0240002
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-371-2600
; TELEFAX: 202-371-2540
; INFORMATION FOR SEQ ID NO: 15:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1835 amino acids
; TYPE: amino acid
; STRANDEDNESS: not relevant
; TOPOLOGY: not relevant
; MOLECULE TYPE: protein
; US-08-836-325-15

Query Match 85.7%; Score 24; DB 3; Length 1835;
Best Local Similarity 100.0%; Pred. No. 3.1e-16;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 5 FVSYIIISFLVVMNYIAVILENF 28
1604 FVSYIIISFLVVMNYIAVILENF 1627

RESULT 13
US-09-457-571-15
; Sequence 15, Application US/09457571
; Patent No. 6703486
; GENERAL INFORMATION:
; APPLICANT: Mandel, Gail
; APPLICANT: Halegoua, Simon
; TITLE OF INVENTION: Peripheral Nervous System Specific
; TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
; TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
; TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
; TITLE OF INVENTION: thereof
; NUMBER OF SEQUENCES: 19
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C
; STREET: 1100 New York Ave., N. W., Suite 600
; CITY: Washington
; STATE: DC


```

COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/457,571
FILING DATE: 09-DEC-1999
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/836,325
FILING DATE: 02-MAY-1997
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917.0240003
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 15:
SEQUENCE CHARACTERISTICS:
LENGTH: 1835 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: not relevant
MOLECULE TYPE: protein
US-09-457-571-15

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Query Match      85.7%; Score 24; DB 4; Length 1835;
Best Local Similarity 100.0%; Pred. No. 3.1e-16;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Cy 5 FVSYSIIISFLVVMNYIAVILENF 28
Db 1604 FVSYSIIISFLVVMNYIAVILENF 1627

RESULT 14
US-08-836-325-16
Sequence 16, Application US/08836325
Patent No. 6110672
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Halegoua, Simon
APPLICANT: Borden, Laurence A.
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Thereof, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
TITLE OF INVENTION: Thereof
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSER: STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS

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SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/836,325
FILING DATE: 2-MAY-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917.0240002
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 16:
SEQUENCE CHARACTERISTICS:
LENGTH: 1969 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-836-325-16

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Query Match      85.7%; Score 24; DB 3; Length 1969;
Best Local Similarity 100.0%; Pred. No. 3.3e-16;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Cy 5 FVSYSIIISFLVVMNYIAVILENF 28
Db 1721 FVSYSIIISFLVVMNYIAVILENF 1744

RESULT 15
US-09-457-571-16
Sequence 16, Application US/09457571
Patent No. 6703486
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Halegoua, Simon
APPLICANT: Borden, Laurence A.
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Thereof, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
TITLE OF INVENTION: Thereof
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSER: STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N. W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/457,571
FILING DATE: 09-DEC-1999
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/836,325
FILING DATE: 02-MAY-1997
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251

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; FILING DATE: 02-NOV-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/482,401
; FILING DATE: 07-JUN-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/334,029
; FILING DATE: 02-NOV-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Ludwig, Steven R.
; REGISTRATION NUMBER: 36,203
; REFERENCE/DOCKET NUMBER: 0917.0240003
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-371-2600
; TELEFAX: 202-371-2540
; INFORMATION FOR SEQ ID NO: 16:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1969 amino acids
; TYPE: amino acid
; STRANDEDNESS: not relevant
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-09-457-571-16
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Query Match      85.7%; Score 24; DB 4; Length 1969;
Best Local Similarity 100.0%; Pred. No. 3.3e-16;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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Qy 5 FVSYYIIISFLVVMNYIAVILLENF 28

Db 1721 FVSYYIIISFLVVMNYIAVILLENF 1744

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RESULT 16
US-09-976-594-757
; Sequence 757, Application US/0976594
; Patent No. 6673549
; GENERAL INFORMATION:
; APPLICANT: Furness, Michael
; APPLICANT: Buchbinder, Jenny
; TITLE OF INVENTION: GENES EXPRESSED IN C3A LIVER CELL CULTURES TREATED WITH STEROIDS
; FILE REFERENCE: PA-0041 US
; CURRENT APPLICATION NUMBER: US/09/976,594
; CURRENT FILING DATE: 2001-10-12
; PRIOR APPLICATION NUMBER: 60/240,409
; PRIOR FILING DATE: 2000-10-12
; NUMBER OF SEQ ID NOS: 1143
; SOFTWARE: PERL Program
; SEQ ID NO 757
; LENGTH: 1977
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; OTHER INFORMATION: Incyte ID No. 6673549 1719478CD1
US-09-976-594-757
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Query Match      85.7%; Score 24; DB 4; Length 1977;
Best Local Similarity 100.0%; Pred. No. 3.3e-16;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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Qy 5 FVSYYIIISFLVVMNYIAVILLENF 28

Db 1729 FVSYYIIISFLVVMNYIAVILLENF 1752

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RESULT 17
US-09-919-039-367
; Sequence 367, Application US/09919039
; Patent No. 6727066
; GENERAL INFORMATION:
; APPLICANT: Kaseer, Matthew R.
; TITLE OF INVENTION: GENES EXPRESSED IN TREATED HUMAN C3A LIVER CELL CULTURES
; FILE REFERENCE: PA-0035 US
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; CURRENT APPLICATION NUMBER: US/09/919,039
; CURRENT FILING DATE: 2002-09-09
; PRIOR APPLICATION NUMBER: 60/222,113
; PRIOR FILING DATE: 2000-07-28
; NUMBER OF SEQ ID NOS: 401
; SOFTWARE: PERL Program
; SEQ ID NO 367
; LENGTH: 1977
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; OTHER INFORMATION: Incyte ID No. 6727066 1719478CD1
US-09-919-039-367
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Query Match      85.7%; Score 24; DB 4; Length 1977;
Best Local Similarity 100.0%; Pred. No. 3.3e-16;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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Qy 5 FVSYYIIISFLVVMNYIAVILLENF 28

Db 1729 FVSYYIIISFLVVMNYIAVILLENF 1752

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RESULT 18
US-08-836-325-10
; Sequence 10, Application US/08836325
; Patent No. 6110672
; GENERAL INFORMATION:
; APPLICANT: Mandel, Gal1
; APPLICANT: Halegoua, Simon
; APPLICANT: Borden, Laurence A.
; TITLE OF INVENTION: Peripheral Nervous System Specific
; TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
; TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
; TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
; TITLE OF INVENTION: Thereof
; NUMBER OF SEQUENCES: 19
; CORRESPONDENCE ADDRESS:
; ADDRESSER: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
; STREET: 1100 New York Ave., N. W., Suite 600
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20005-3934
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/836,325
; FILING DATE: 2-MAY-1997
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/14251
; FILING DATE: 02-NOV-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/482,401
; FILING DATE: 07-JUN-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/334,029
; FILING DATE: 02-NOV-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Ludwig, Steven R.
; REGISTRATION NUMBER: 36,203
; REFERENCE/DOCKET NUMBER: 0917.0240002
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-371-2540
; TELEFAX: 202-371-2540
; INFORMATION FOR SEQ ID NO: 10:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1984 amino acids
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TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-836-325-10

Query Match 85.7%; Score 24; DB 3; Length 1984;
Best Local Similarity 100.0%; Pred. No. 3.3e-16;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 FVSYYIIISFLVVMNYIAVILENF 28
DB 1738 FVSYYIIISFLVVMNYIAVILENF 1761

RESULT 19
US-09-457-571-10
Sequence 10, Application US/09457571
Patent No. 6703486
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Halogoua, Simon
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
TITLE OF INVENTION: Thereof
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N.W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/457,571
FILING DATE: 09-DEC-1999
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/836,325
FILING DATE: 02-MAY-1997
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917,0240003
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 10:
SEQUENCE CHARACTERISTICS:
LENGTH: 1984 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-09-457-571-10

Query Match 85.7%; Score 24; DB 4; Length 1984;
Best Local Similarity 100.0%; Pred. No. 3.3e-16;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 FVSYYIIISFLVVMNYIAVILENF 28
DB 1738 FVSYYIIISFLVVMNYIAVILENF 1761

RESULT 20
US-08-836-325-11
Sequence 11, Application US/08836325
Patent No. 6110672
GENERAL INFORMATION:
APPLICANT: Mandel, Gail
APPLICANT: Halogoua, Simon
APPLICANT: Borden, Laurence A.
TITLE OF INVENTION: Peripheral Nervous System Specific
TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
TITLE OF INVENTION: Thereof
NUMBER OF SEQUENCES: 19
CORRESPONDENCE ADDRESS:
ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C
STREET: 1100 New York Ave., N.W., Suite 600
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/836,325
FILING DATE: 2-MAY-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917,0240002
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 11:
SEQUENCE CHARACTERISTICS:
LENGTH: 1989 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: not relevant
MOLECULE TYPE: protein
US-08-836-325-11

Query Match 85.7%; Score 24; DB 3; Length 1989;
Best Local Similarity 100.0%; Pred. No. 3.3e-16;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 FVSYYIIISFLVVMNYIAVILENF 28
DB 1740 FVSYYIIISFLVVMNYIAVILENF 1763

RESULT 21
US-08-836-325-12
Sequence 12, Application US/08836325

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; Patent No. 6110672
; GENERAL INFORMATION:
; APPLICANT: Mandel, Gail
; APPLICANT: Halegoua, Simon
; APPLICANT: Borden, Laurence A.
; TITLE OF INVENTION: Peripheral Nervous System Specific
; TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
; TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
; TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
; TITLE OF INVENTION: Thereof
; NUMBER OF SEQUENCES: 19
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C.
; STREET: 1100 New York Ave., N. W., Suite 600
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20005-3934
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/836,325
; FILING DATE: 2-MAY-1997
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/14251
; FILING DATE: 02-NOV-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/482,401
; FILING DATE: 07-JUN-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/334,029
; FILING DATE: 02-NOV-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Ludwig, Steven R.
; REGISTRATION NUMBER: 36,203
; REFERENCE/DOCKET NUMBER: 0917.0240002
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-371-2600
; TELEFAX: 202-371-2540
; INFORMATION FOR SEQ ID NO: 12:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1989 amino acids
; TYPE: amino acid
; STRANDEDNESS: not relevant
; TOPOLOGY: not relevant
; MOLECULE TYPE: protein
; US-08-836-325-12

Query Match      85.7%; Score 24; DB 3; Length 1989;
Best Local Similarity 100.0%; Pred. No. 3.3e-16;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      5 FVSYYIIISFLVVVMYIAVILLENF 28
DB      1740 FVSYYIIISFLVVVMYIAVILLENF 1763

RESULT 22
US-09-457-571-11
; Sequence 11, Application US/09457571
; Patent No. 6703486
; GENERAL INFORMATION:
; APPLICANT: Mandel, Gail
; APPLICANT: Halegoua, Simon
; TITLE OF INVENTION: Peripheral Nervous System Specific
; TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
; TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
; TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
; TITLE OF INVENTION: Thereof

```

```

; NUMBER OF SEQUENCES: 19
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C.
; STREET: 1100 New York Ave., N. W., Suite 600
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20005-3934
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/457,571
; FILING DATE: 09-DEC-1999
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/836,325
; FILING DATE: 02-MAY-1997
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/14251
; FILING DATE: 02-NOV-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/482,401
; FILING DATE: 07-JUN-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/334,029
; FILING DATE: 02-NOV-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Ludwig, Steven R.
; REGISTRATION NUMBER: 36,203
; REFERENCE/DOCKET NUMBER: 0917.0240003
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-371-2600
; TELEFAX: 202-371-2540
; INFORMATION FOR SEQ ID NO: 11:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1989 amino acids
; TYPE: amino acid
; STRANDEDNESS: not relevant
; TOPOLOGY: not relevant
; MOLECULE TYPE: protein
; US-09-457-571-11

Query Match      85.7%; Score 24; DB 4; Length 1989;
Best Local Similarity 100.0%; Pred. No. 3.3e-16;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      5 FVSYYIIISFLVVVMYIAVILLENF 28
DB      1740 FVSYYIIISFLVVVMYIAVILLENF 1763

RESULT 23
US-09-457-571-12
; Sequence 12, Application US/09457571
; Patent No. 6703486
; GENERAL INFORMATION:
; APPLICANT: Mandel, Gail
; APPLICANT: Halegoua, Simon
; TITLE OF INVENTION: Peripheral Nervous System Specific
; TITLE OF INVENTION: Sodium Channels, DNA Encoding Therefor, Crystallization,
; TITLE OF INVENTION: X-ray Diffraction, Computer Molecular Modeling, Rational
; TITLE OF INVENTION: Drug Design, Drug Screening, and Methods of Making and Using
; TITLE OF INVENTION: Thereof
; NUMBER OF SEQUENCES: 19
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: STERN, KESSLER, GOLDSTEIN & FOX P.L.L.C.
; STREET: 1100 New York Ave., N. W., Suite 600
; CITY: Washington
; STATE: DC
; COUNTRY: USA

```

ZIP: 20005-3934
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/457,571
FILING DATE: 09-DEC-1999
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/836,325
FILING DATE: 02-MAY-1997
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/US95/14251
FILING DATE: 02-NOV-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/482,401
FILING DATE: 07-JUN-1995
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/334,029
FILING DATE: 02-NOV-1994
ATTORNEY/AGENT INFORMATION:
NAME: Ludwig, Steven R.
REGISTRATION NUMBER: 36,203
REFERENCE/DOCKET NUMBER: 0917,0240003
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-371-2600
TELEFAX: 202-371-2540
INFORMATION FOR SEQ ID NO: 12:
SEQUENCE CHARACTERISTICS:
LENGTH: 1989 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: not relevant
MOLECULE TYPE: protein
US-09-457-571-12

Query Match 85.7%; Score 24; DB 4; Length 1989;
Best Local Similarity 100.0%; Pred. No. 3.3e-16;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 FVSIIISFLVVMYIAVLENF 28
DB 1740 FVSIIISFLVVMYIAVLENF 1763

RESULT 24
US-08-605-284B-16
Sequence 16, Application US/08605284B
Patent No. 6060271
GENERAL INFORMATION:
APPLICANT: MALEWSKI, JOSE L.
REGISTRATION NUMBER: 34,103
REFERENCE/DOCKET NUMBER: 19603/800 (CRF D-1705)
TITLE OF INVENTION: VOLTAGE GATED SODIUM CHANNELS FROM
NUMBER OF SEQUENCES: 23
CORRESPONDENCE ADDRESS:
ADDRESSEE: NIXON, HARGRAVE, DEVANS & DOYLE LLP
STREET: CLINTON SQUARE, P.O. BOX 1051
CITY: ROCHESTER
STATE: NEW YORK
COUNTRY: USA
ZIP: 14603
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/605,284B
FILING DATE: 09-FEB-1996
CLASSIFICATION: 424

ATTORNEY/AGENT INFORMATION:
NAME: BRAMAN, SUSAN J.
REGISTRATION NUMBER: 34,103
REFERENCE/DOCKET NUMBER: 19603/800 (CRF D-1705)
TELECOMMUNICATION INFORMATION:
TELEPHONE: 716-263-1636
TELEFAX: 716-263-1600
INFORMATION FOR SEQ ID NO: 16:
SEQUENCE CHARACTERISTICS:
LENGTH: 310 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-605-284B-16

Query Match 60.7%; Score 17; DB 3; Length 310;
Best Local Similarity 100.0%; Pred. No. 7.9e-10;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 12 SFLVVMYIAVLENF 28
DB 294 SFLVVMYIAVLENF 310

RESULT 25
US-08-605-284B-10
Sequence 10, Application US/08605284B
Patent No. 6060271
GENERAL INFORMATION:
APPLICANT: MALEWSKI, JOSE L.
REGISTRATION NUMBER: 34,103
REFERENCE/DOCKET NUMBER: 19603/800 (CRF D-1705)
TITLE OF INVENTION: VOLTAGE GATED SODIUM CHANNELS FROM
NUMBER OF SEQUENCES: 23
CORRESPONDENCE ADDRESS:
ADDRESSEE: NIXON, HARGRAVE, DEVANS & DOYLE LLP
STREET: CLINTON SQUARE, P.O. BOX 1051
CITY: ROCHESTER
STATE: NEW YORK
COUNTRY: USA
ZIP: 14603
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/605,284B
FILING DATE: 09-FEB-1996
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: BRAMAN, SUSAN J.
REGISTRATION NUMBER: 34,103
REFERENCE/DOCKET NUMBER: 19603/800 (CRF D-1705)
TELECOMMUNICATION INFORMATION:
TELEPHONE: 716-263-1636
TELEFAX: 716-263-1600
INFORMATION FOR SEQ ID NO: 10:
SEQUENCE CHARACTERISTICS:
LENGTH: 310 amino acids
TYPE: amino acid
STRANDEDNESS: not relevant
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-605-284B-10

Query Match 57.1%; Score 16; DB 3; Length 310;
Best Local Similarity 100.0%; Pred. No. 8.2e-09;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GIFFVSYIIISFLVV 16
DB 1 GIFFVSYIIISFLVV 16

Fri Jan 28 09:32:03 2005

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Page 12

Db 283 GIFFVSYIISFLV 298

Search completed: January 27, 2005, 17:54:20
Job time : 23.5 secs